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(54) Title: BICYCLIC HETEROAROMATIC COMPOUNDS AS KINASE INHIBITORS

(57) Abstract: A series of 5-6 fused ring bicyclic heteroaromatic derivatives, based in particular on the 5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine ring system, being inhibitors of p38 kinase, are accordingly of use in medicine, for example in the treatment and/or prevention of immune or inflammatory disorders.

BICYCLIC HETEROAROMATIC COMPOUNDS AS KINASE INHIBITORS

5 This invention relates to a series of 5-6 fused ring bicyclic heteroaromatic derivatives, to compositions containing them, to processes for their preparation and to their use in medicine.

Immune and inflammatory responses involve a variety of cell types with control and co-ordination of the various interactions occurring *via* both cell-cell contacts (e.g. 10 integrin interactions with their receptors) and by way of intercellular signalling molecules. A large number of different signalling molecules are involved including cytokines, lymphocytes, chemokines and growth factors.

Cells respond to such intercellular signalling molecules by means of intracellular signalling mechanisms that include protein kinases, phosphatases and phospholipases. 15 There are five classes of protein kinase of which the major ones are the tyrosine kinases and the serine/threonine kinases [Hunter, T., *Methods in Enzymology (Protein Kinase Classification)*, p. 3, Hunter, T. and Sefton, B.M. eds., vol. 200, Academic Press, San Diego, 1991].

One sub-class of serine/threonine kinases is the mitogen activating protein (MAP) 20 kinases of which there are at least three families which differ in the sequence and size of the activation loop [Adams, J. L. et al., *Progress in Medicinal Chemistry*, pp. 1-60, King, F.D. and Oxford, A.W. eds., vol. 38, Elsevier Science, 2001]: the extracellular regulated kinases (ERKs); the c-Jun NH₂ terminal kinases or stress activated kinases (JNKs or SAP kinases); and the p38 kinases, which have a threonine-glycine-tyrosine (TGY) activation 25 motif. Both the JNKs and p38 MAP kinases are primarily activated by stress stimuli including, but not limited to, proinflammatory cytokines, e.g. tumour necrosis factor (TNF) and interleukin-1 (IL-1), ultraviolet light, endotoxin and chemical or osmotic shock.

Four isoforms of p38 have been described (p38 $\alpha/\beta/\gamma/\delta$). The human p38 α 30 enzyme was initially identified as a target of cytokine-suppressive anti-inflammatory drugs (CSAIDs) and the two isoenzymes found were initially termed CSAID binding protein-1 and -2 (CSBP-1 and CSBP-2 respectively) [Lee, J. C. et al., *Nature (London)*, 1994, 372, 739-46]. CSBP-2 is now widely referred to as p38 α and differs from CSBP-1

in an internal sequence of 25 amino acids as a result of differential splicing of two exons that are conserved in both mouse and human [McDonnell, P.C. *et al.*, *Genomics*, 1995, 29, 301-2]. CSBP-1 and p38 α are expressed ubiquitously and there is no difference between the two isoforms with respect to tissue distribution, activation profile, substrate preference or CSAID binding. A second isoform is p38 β which has 70% identity with p38 α . A second form of p38 β termed p38 β 2 is also known and of the two this is believed to be the major form. p38 α and p38 β 2 are expressed in many different tissues. However, in monocytes and macrophages p38 α is the predominant kinase activity [Lee, J.C., *ibid*; Jing, Y. *et al.*, *J. Biol. Chem.*, 1996, 271, 10531-34; Hale, K.K. *et al.*, *J. Immun.*, 1999, 162, 4246-52]. p38 γ and p38 δ (also termed SAP kinase-3 and SAP kinase-4 respectively) have ~63% and ~61% homology to p38 α respectively. p38 γ is predominantly expressed in skeletal muscle whilst p38 δ is found in testes, pancreas, prostate, small intestine and in certain endocrine tissues.

All p38 homologues and splice variants contain a 12 amino acid activation loop that includes a Thr-Gly-Tyr motif. Dual phosphorylation of both Thr-180 and Tyr-182 in the TGY motif by a dual specificity upstream kinase is essential for the activation of p38 and results in a >1000-fold increase in specific activity of these enzymes [Doza, Y.N. *et al.*, *FEBS Lett.*, 1995, 364, 7095-8012]. This dual phosphorylation is effected by MKK6 and, under certain conditions, the related enzyme MKK3 [Enslen, H. *et al.*, *J. Biol. Chem.*, 1998, 273, 1741-48]. MKK3 and MKK6 belong to a family of enzymes termed MAPKK (mitogen activating protein kinase kinase) which are in turn activated by MAPKKK (mitogen activating kinase kinase kinase) otherwise known as MAP3K.

Several MAP3Ks have been identified that are activated by a wide variety of stimuli including environmental stress, inflammatory cytokines and other factors. MEKK4/MTK1 (MAP or ERK kinase kinase/MAP three kinase-1), ASK1 (apoptosis stimulated kinase) and TAK1 (TGF- β -activated kinase) are some of the enzymes identified as upstream activators of MAPKKs. MEKK4/MTK1 is thought to be activated by several GADD-45-like genes that are induced in response to environmental stimuli and which eventually lead to p38 activation [Takekawa, M. and Saito, H., *Cell*, 1998, 95, 521-30]. TAK1 has been shown to activate MKK6 in response to transforming growth factor- β (TGF- β). TNF-stimulated activation of p38 is believed to be mediated by the recruitment of TRAF2 (TNF receptor associated factor) and the Fas adaptor protein, Daxx, which results in the activation of ASK1 and subsequently p38.

Several substrates of p38 have been identified including other kinases [e.g. MAPK activated protein kinase 2/3/5 (MAPKAP 2/3/5), p38 regulated/activated protein kinase (PRAK), MAP kinase-interacting kinase 1/2 (MNK1/2), mitogen- and stress-activated protein kinase 1 (MSK1/RLPK) and ribosomal S6 kinase-B (RSK-B)], transcription factors [e.g. activating transcription factor 2/6 (ATF2/6), monocyte-enhancer factor-2A/C (MEF2A/C), C/EBP homologous protein (CHOP), Elk1 and Sap-1a1] and other substrates [e.g. cPLA₂, p47phox].

MAPKAP K2 is activated by p38 in response to environmental stress. Mice engineered to lack MAPKAP K2 do not produce TNF in response to lipopolysaccharide (LPS). Production of several other cytokines such as IL-1, IL-6, IFN- γ and IL-10 is also partially inhibited [Kotlyarov, A. et al., *Nature Cell Biol.*, 1999, 1, 94-7]. Further, MAPKAP K2 from embryonic stem cells from p38 α null mice was not activated in response to stress and these cells did not produce IL-6 in response to IL-1 [Allen, M. et al., *J. Exp. Med.*, 2000, 191, 859-69]. These results indicate that MAPKAP K2 is not only essential for TNF and IL-1 production but also for signalling induced by cytokines. In addition, MAPKAP K2/3 phosphorylate and thus regulate heat shock proteins HSP 25 and HSP 27 which are involved in cytoskeletal reorganization.

Several small molecule inhibitors of p38 have been reported which inhibit IL-1 and TNF synthesis in human monocytes at concentrations in the low μ M range [Lee, J.C. et al., *Int. J. Immunopharm.*, 1988, 10, 835] and exhibit activity in animal models which are refractory to cyclooxygenase inhibitors [Lee, J.C. et al., *Annals N.Y. Acad. Sci.*, 1993, 696, 149]. In addition, these small molecule inhibitors are known to decrease the synthesis of a wide variety of pro-inflammatory proteins including IL-6, IL-8, granulocyte/macrophage colony-stimulating factor (GM-CSF) and cyclooxygenase-2 (COX-2). TNF-induced phosphorylation and activation of cytosolic PLA₂, TNF-induced expression of VCAM-1 on endothelial cells and IL-1 stimulated synthesis of collagenase and stromelysin are also inhibited by such small molecule inhibitors of p38 [Cohen, P., *Trends Cell Biol.*, 1997, 7, 353-61].

A variety of cells including monocytes and macrophages produce TNF and IL-1. Excessive or unregulated TNF production is implicated in a number of disease states including Crohn's disease, ulcerative colitis, pyresis, rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, gouty arthritis and other arthritic conditions, toxic shock syndrome, endotoxic shock, sepsis, septic shock, gram negative sepsis, bone resorption

diseases, reperfusion injury, graft vs. host reaction, allograft rejection, adult respiratory distress syndrome, chronic pulmonary inflammatory disease, silicosis, pulmonary sarcoidosis, cerebral malaria, scar tissue formation, keloid formation, fever and myalgias due to infection, such as influenza, cachexia secondary to acquired immune deficiency

5 syndrome (AIDS), cachexia secondary to infection or malignancy, AIDS or AIDS related complex.

Excessive or unregulated IL-1 production has been implicated in rheumatoid arthritis, osteoarthritis, traumatic arthritis, rubella arthritis, acute synovitis, psoriatic arthritis, cachexia, Reiter's syndrome, endotoxemia, toxic shock syndrome, tuberculosis,

10 atherosclerosis, muscle degeneration, and other acute or chronic inflammatory diseases such as the inflammatory reaction induced by endotoxin or inflammatory bowel disease.

In addition, IL-1 has been linked to diabetes and pancreatic β cells [Dinarello, C.A., *J. Clinical Immunology*, 1985, 5, 287-97].

IL-8 is a chemotactic factor produced by various cell types including endothelial 15 cells, mononuclear cells, fibroblasts and keratinocytes. IL-1, TNF and LPS all induce the production of IL-8 by endothelial cells. *In vitro* IL-8 has been shown to have a number of functions including being a chemoattractant for neutrophils, T-lymphocytes and basophils. IL-8 has also been shown to increase the surface expression of Mac-1 (CD11b/CD18) on neutrophils without *de novo* protein synthesis, which may contribute 20 to increased adhesion of neutrophils to vascular endothelial cells. Many diseases are characterised by massive neutrophil infiltration. Histamine release from basophils (in both atopic and normal individuals) is induced by IL-8 as is lysozomal enzyme release and respiratory burst from neutrophils.

The central role of IL-1 and TNF together with other leukocyte-derived cytokines 25 as important and critical inflammatory mediators is well documented. The inhibition of these cytokines has been shown or would be expected to be of benefit in controlling, alleviating or reducing many of these disease states.

The central position that p38 occupies within the cascade of signalling molecules mediating extracellular to intracellular signalling, and its influence over not only IL-1, 30 TNF and IL-8 production but also the synthesis and/or action of other pro-inflammatory proteins (e.g. IL-6, GM-CSF, COX-2, collagenase and stromelysin), make it an attractive target for inhibition by small molecule inhibitors with the expectation that such inhibition would be a highly effective mechanism for regulating the excessive and destructive

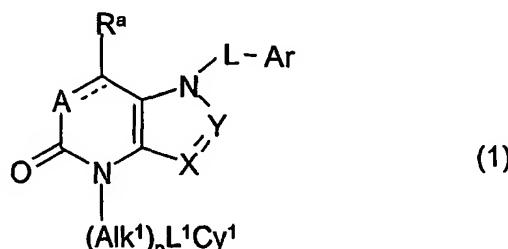
activation of the immune system. Such an expectation is supported by the potent and diverse anti-inflammatory activities described for p38 kinase inhibitors [Adams, *ibid*; Badger *et al.*, *J. Pharmacol. Exp. Ther.*, 1996, **279**, 1453-61; Griswold *et al.*, *Pharmacol. Commun.*, 1996, **7**, 323-29].

5 Certain *N,N'*-dibenzylated 2-purinones have been described (Andresen *et al.*, *Tetrahedron*, **52**, 12979-12992).

The present invention provides a class of compounds which are potent and selective inhibitors of p38 kinase, especially p38 α , p38 β and p38 β 2, and splice variants thereof. The compounds in accordance with the present invention are thus of use in
10 medicine, for example in the prophylaxis and treatment of immune or inflammatory disorders.

In addition, the compounds according to the present invention may be used as pharmacological standards for use in the development of new biological tests and in the search for new pharmacological agents. Thus, the compounds according to this invention
15 may be useful as radioligands in assays for detecting compounds capable of binding to the human p38 enzyme.

Thus according to one aspect of the invention we provide a compound of formula (1):



20

wherein:

the dashed line joining A and C(R^a) is present and represents a bond and A is a -N= atom or a -C(R^b)= group, or the dashed line is absent and A is a -C(R^b)(R^c)- or
25 -N(R^d)- group;

R^a, R^b and R^c is each independently a hydrogen or halogen atom or an optionally substituted alkyl, -CN, -CO₂R¹ (where R¹ is a hydrogen atom or an optionally substituted alkyl group) or -CONR¹R² group (where R² is a hydrogen atom or an optionally substituted alkyl group);

R^d is a hydrogen atom or an alkyl group;

X and Y is each a nitrogen atom or a $-C(R^e)=$ or $-C(Alk^2R^e)=$ group [where Alk^2 is an optionally substituted alkylene, alkenylene or alkynylene chain and R^e is a hydrogen or halogen atom or a $-CN$, $-OR^1$, $-CO_2R^1$, $-C(X^a)R^1$ (where X^a is an oxygen or sulphur atom),

- 5 $-Cy^2$ (where Cy^2 is an optionally substituted, saturated or unsaturated non-aromatic carbocyclic ring optionally containing one or more $-O-$, $-S-$, $-NH-$ or $-C(X^a)-$ atoms or groups), $-NR^{1a}R^{2a}$ (where R^{1a} and R^{2a} , which may be the same or different, is each a hydrogen atom or an optionally substituted alkyl or Cy^2 group, or together with the nitrogen atom to which they are attached form an optionally substituted, saturated or unsaturated cyclicamino ring optionally containing one or more $-O-$ or $-S-$ atoms or $-NH-$ or $-C(X^a)-$ groups), $-C(X^a)NR^{1a}R^{2a}$, $-S(O)_2NR^{1a}R^{2a}$, $-N(R^{3a})C(X^a)R^1$ (where R^{3a} is a hydrogen atom or an optionally substituted alkyl group), $-N(R^{3a})C(X^a)NR^{1a}R^{2a}$, $-N(R^{3a})S(O)_2R^1$, $-N[S(O)_2R^1]_2$, $-N(R^{3a})S(O)_2NR^{1a}R^{2a}$, $-N(R^{3a})C(O)OR^1$, $-N(R^{3a})C(NR^1)NR^{1a}R^{2a}$, $-C(R^1)NOR^2$, $-C(NR^1)NR^{1a}R^{2a}$, $-C(X^a)NR^{1a}OR^{2a}$ or $-C(O)N(R^{3a})NR^{1a}R^{2a}$ group];
- 10
- 15 L is a $-C(O)-$, $-C(S)-$ or $-C(R^{1f})(R^{1g})-$ group (where R^{1f} and R^{1g} , which may be the same or different, is each a hydrogen atom or a straight or branched C_{1-3} alkyl group optionally substituted by one, two or three fluorine atoms, or R^{1f} and R^{1g} together with the carbon atom to which they are attached form a cyclopropyl group), or a $-CH_2CH_2-$ group;
- n is zero or the integer 1;
- 20 Alk¹ is an optionally substituted aliphatic or heteroaliphatic chain;
- L¹ is a covalent bond or a linker atom or group;
- Cy¹ is an optionally substituted cycloaliphatic, polycycloaliphatic, heterocycloaliphatic, polyheterocycloaliphatic, aromatic or heteroaromatic group, or is additionally a hydrogen atom when n is the integer 1 and/or L¹ is a linker atom or group;
- 25 and
- Ar is an optionally substituted aromatic or heteroaromatic group;
- provided that the compound of formula (1) is other than 3,7-dibenzyl-3,7-dihydro-2H-purinone;
- and the salts, solvates, hydrates and N-oxides thereof.
- 30 The present invention also provides a compound of formula (1) as depicted above, or a salt, solvate, hydrate or N-oxide thereof, wherein L is a $-C(O)-$, $-C(S)-$ or $-C(R^{1f})(R^{1g})-$ group, and the remaining substituents are as defined above.

It will be appreciated that compounds of formula (1) may have one or more chiral centres, and exist as enantiomers or diastereomers. The invention is to be understood to extend to all such enantiomers, diastereomers and mixtures thereof, including racemates. Formula (1) and the formulae hereinafter are intended to represent all individual isomers

5 and mixtures thereof, unless stated or shown otherwise. In addition, compounds of formula (1) may exist as tautomers, for example keto ($\text{CH}_2\text{C}=\text{O}$)-enol ($\text{CH}=\text{CHOH}$) tautomers. Formula (1) and the formulae hereinafter are intended to represent all individual tautomers and mixtures thereof, unless stated otherwise.

The following general terms as used herein in relation to compounds of the

10 invention and intermediates thereto have the stated meaning below unless specifically defined otherwise.

Thus, as used herein, the term "alkyl" whether present as a group or part of a group includes straight or branched C_{1-6} alkyl groups, for example C_{1-4} alkyl groups such as methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, *sec*-butyl, isobutyl or *tert*-butyl groups.

15 Similarly, the terms "alkenyl" and "alkynyl" are intended to mean straight or branched C_{2-6} alkenyl or C_{2-6} alkynyl groups such as C_{2-4} alkenyl or C_{2-4} alkynyl groups.

Optional substituents which may be present on any of these alkyl, alkenyl or alkynyl groups include one, two, three or more substituents where each substituent may be the same or different and is selected from a halogen atom and an -OH, -CO₂H, -CO₂R⁴

20 (where R⁴ is an optionally substituted straight or branched C_{1-6} alkyl group), -CONH₂, -CONHR⁴, -CON(R⁴)₂, -COR⁴, C_{1-6} alkoxy, halo(C_{1-6} alkoxy), -S(O)R⁴, -S(O)₂R⁴, C_{1-6} alkylthio, -NH₂, -NHR⁴, -N(R⁴)₂, -S(O)₂NH₂, -S(O)₂NHR⁴, -S(O)₂N(R⁴)₂, -NHC(O)R⁴, -N(R⁴)C(O)R⁴, -NHS(O)₂R⁴, -N(R⁴)S(O)₂R⁴, -NHS(O)₂NH₂, -NHS(O)₂NHR⁴, -NHS(O)₂N(R⁴)₂, -N(R⁴)S(O)₂NH₂, -N(R⁴)S(O)₂NHR⁴ or -N(R⁴)S(O)₂N(R⁴)₂ group.

25 Where two R⁴ groups are present in a group, these may be the same or different or may be joined, together with the N atom to which they are attached, to form a heterocyclic ring. Such heterocyclic rings may be optionally interrupted by a further heteroatom or heteroatom-containing group selected from -O-, -S-, -NH-, -N(R⁴)-, -C(O)- and -C(S)- groups. Particular examples of such heterocyclic rings include piperidinyl, pyrazolidinyl,

30 morpholinyl, thiomorpholinyl, pyrrolidinyl, imidazolidinyl or piperazinyl rings.

The term "halogen" as used herein is intended to include fluorine, chlorine, bromine and iodine atoms.

The term "haloalkyl" is intended to include those alkyl groups particularly mentioned above substituted by one, two or three of the halogen atoms just described. Particular examples of such groups include -CF₃, -CCl₃, -CHF₂, -CHCl₂, -CH₂F and -CH₂Cl groups. Each haloalkyl group may be optionally substituted by one or more

5 hydroxy groups and the term is to be understood to include such substituted groups, such as -C(OH)CF₃)₂ groups.

The term "alkoxy" as used herein is intended to include straight or branched C₁₋₆ alkoxy, e.g. C₁₋₄ alkoxy such as methoxy, ethoxy, *n*-propoxy, isopropoxy, *n*-butoxy, *sec*-butoxy, isobutoxy and *tert*-butoxy. "Haloalkoxy" as used herein includes any of these

10 alkoxy groups substituted by one, two or three halogen atoms as described above. Particular examples include -OCF₃, -OCCl₃, -OCHF₂, -OCHCl₂, -OCH₂F and -OCH₂Cl groups.

As used herein, the term "alkylthio" is intended to include straight or branched C₁₋₆ alkylthio, e.g. C₁₋₄ alkylthio such as methylthio or ethylthio.

15 When Alk¹ is present in compounds of formula (1) as an optionally substituted aliphatic chain it may be an optionally substituted C₁₋₁₀ aliphatic chain. Particular examples include optionally substituted straight or branched chain C₁₋₆ alkylene, C₂₋₆ alkenylene and C₂₋₆ alkynylene chains.

Particular examples of aliphatic chains represented by Alk¹ include optionally

20 substituted -CH₂-, -CH₂CH₂-, -CH(CH₃)CH₂-, -(CH₂)₂CH₂-, -(CH₂)₃CH₂-, -CH(CH₃)(CH₂)₂CH₂-, -CH₂CH(CH₃)CH₂-, -C(CH₃)₂CH₂-, -CH₂C(CH₃)₂CH₂-, -(CH₂)₂CH(CH₃)CH₂-, -CH(CH₃)CH₂CH₂-, -CH(CH₃)CH₂CH(CH₃)CH₂-, -CH₂CH(CH₃)CH₂CH₂-, -(CH₂)₂C(CH₃)₂CH₂-, -(CH₂)₄CH₂-, -(CH₂)₅CH₂-, -CH=CH-, -CH=CHCH₂-, -CH₂CH=CH-, -CH=CHCH₂CH₂-, -CH₂CH=CHCH₂-, -(CH₂)₂CH=CH-,

25 -C≡C-, -C≡CCH₂-, -CH₂C≡C-, -C≡CCH₂CH₂-, -CH₂C≡CCH₂- and -(CH₂)₂C≡C- chains.

Heteroaliphatic chains represented by Alk¹ in the compounds of formula (1) include the aliphatic chains just described but with each additionally containing one, two, three or four heteroatoms or heteroatom-containing groups. Particular heteroatoms or groups include atoms or groups L² where L² is a linker atom or group. Each L² atom or group may interrupt

30 the aliphatic group, or may be positioned at its terminal carbon atom to connect the group to an adjoining atom or group. Particular examples include optionally substituted -L²CH₂-, -CH₂L²-, -L²CH(CH₃)-, -CH(CH₃)L²-, -CH₂L²CH₂-, -L²CH₂CH₂-, -L²CH₂CH(CH₃)-, -CH(CH₃)CH₂L²-, -CH₂CH₂L²-, -CH₂L²CH₂CH₂-, -CH₂L²CH₂CH₂L²-, -(CH₂)₂L²CH₂-,

$-(CH_2)_3L^2CH_2$, $-L^2(CH_2)_2CH_2$, $-L^2CH_2CH=CH-$, $-CH=CHCH_2L^2$ and $-(CH_2)_2L^2CH_2CH_2$ - chains.

When L^2 is present in heteroaliphatic chains as a linker atom or group it may be any divalent linking atom or group. Particular examples include -O- or -S- atoms and

5 $-C(O)-$, $-C(O)O-$, $-OC(O)-$, $-C(S)-$, $-S(O)-$, $-S(O)_2-$, $-N(R^3)-$ (where R^3 is a hydrogen atom or a straight or branched alkyl group), $-N(R^3)O-$, $-N(R^3)NH-$, $-CON(R^3)-$, $-OC(O)N(R^3)-$, $-CSN(R^3)-$, $-N(R^3)CO-$, $-N(R^3)C(O)O-$, $-N(R^3)CS-$, $-S(O)_2N(R^3)-$, $-N(R^3)S(O)_2-$, $-N(R^3)CON(R^3)-$, $-N(R^3)CSN(R^3)-$ or $-N(R^3)SO_2N(R^3)-$ groups. Where L^2 contains two R^3 groups these may be the same or different.

10 The optional substituents which may be present on aliphatic or heteroaliphatic chains represented by Alk^1 include one, two or three of those optional substituents described above in relation to the term "alkyl".

When X or Y in compounds of the invention contains an Alk^2 chain, the chain may be an optionally substituted, straight or branched C_{1-6} alkylene, C_{2-6} alkenylene or C_{2-6} alkynylene chain. Particular chains include those just described above for the chain Alk^1 . Where desired, each chain may be substituted by one, two or three substituents as described for Alk^1 .

When a carbocyclic non-aromatic ring Cy^2 is present in compounds of formula (1) it may be an optionally substituted C_{3-7} cycloalkyl, C_{3-7} cycloalkenyl, C_{3-7} heterocycloalkyl or 20 C_{3-7} heterocycloalkenyl ring as generally and particularly described hereinafter in relation to the group Cy^1 . Optional substituents, which may be present on any available carbon or nitrogen atom in the ring, include those described above in relation to the term "alkyl".

When $-NR^{1a}R^{2a}$ is present in compounds of the invention as a cyclicamino ring it may be an optionally substituted heterocycloaliphatic group, as generally and particularly 25 described below in relation to the group Cy^1 , which is attached to the remainder of the molecule through a nitrogen atom. Examples include azetidin-1-yl, pyrrolidin-1-yl, piperidin-1-yl, homopiperidin-1-yl, morpholin-4-yl and piperazin-1-yl, especially pyrrolidin-1-yl. Each ring may be optionally substituted on any available ring carbon or nitrogen atom by a substituent selected from those described above in relation to the term "alkyl".

30 When L^1 is present in compounds of formula (1) as a linker atom or group it may be any such atom or group as hereinbefore described in relation to L^2 linker atoms and groups.

Optionally substituted cycloaliphatic groups represented by the group Cy¹ in compounds of the invention include optionally substituted C₃₋₁₀ cycloaliphatic groups. Particular examples include optionally substituted C₃₋₁₀ cycloalkyl, e.g. C₃₋₇ cycloalkyl, and C₃₋₁₀ cycloalkenyl, e.g. C₃₋₇ cycloalkenyl, groups.

5 Optionally substituted heterocycloaliphatic groups represented by the group Cy¹ include optionally substituted C₃₋₁₀ heterocycloaliphatic groups. Particular examples include optionally substituted C₃₋₁₀ heterocycloalkyl, e.g. C₃₋₇ heterocycloalkyl, and C₃₋₁₀ heterocycloalkenyl, e.g. C₃₋₇ heterocycloalkenyl, groups, each of said groups containing one, two, three or four heteroatoms or heteroatom-containing groups L⁴ in place of or in
10 addition to the ring carbon atoms, where L⁴ is an atom or group as previously defined for L².

15 Optionally substituted polycycloaliphatic groups represented by the group Cy¹ include optionally substituted C₇₋₁₀ bi- or tricycloalkyl and C₇₋₁₀ bi- or tricycloalkenyl groups. Optionally substituted heteropolycycloaliphatic groups represented by the group Cy¹ include optionally substituted C₇₋₁₀ bi- or tricycloalkyl and C₇₋₁₀ bi- or tricycloalkenyl groups containing one, two, three, four or more L⁴ atoms or groups in place of or in addition to the ring carbon atoms.

20 Particular examples of cycloaliphatic, polycycloaliphatic, heterocycloaliphatic and heteropolycycloaliphatic groups represented by the group Cy¹ include optionally substituted cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclobut-2-en-1-yl, cyclopent-2-en-1-yl, cyclopent-3-en-1-yl, adamantyl, norbornyl, norbornenyl, dihydrofuryl, tetrahydrofuryl, tetrahydropyranyl, dihydrothienyl, tetrahydrothienyl, pyrrolinyl, e.g. 2- or 3-pyrrolinyl, pyrrolidinyl, pyrrolidinonyl, oxazolidinyl, oxazolidinonyl, dioxolanyl, e.g. 1,3-dioxolanyl, imidazolinyl, e.g. imidazolin-2-yl, imidazolidinyl, pyrazolinyl, e.g. pyrazolin-2-yl, pyrazolidinyl, 5,6-dihydro-2(1H)-pyrazinonyl, tetrahydropyrimidinyl, thiazolinyl, thiazolidinyl, pyranyl, e.g. 2- or 4-pyranyl, piperidinyl, homopiperidinyl, heptamethyleneiminy, piperidinonyl, 1,4-dioxanyl, morpholinyl, morpholinonyl, 1,4-dithianyl, thiomorpholinyl, piperazinyl, homopiperazinyl, 1,3,5-trithianyl, oxazinyl, e.g. 2H-1,3-, 6H-1,3-, 6H-1,2-, 2H-1,2- or
25 4H-1,4-oxazinyl, 1,2,5-oxathiazinyl, isoxazinyl, e.g. o- or p-isoxazinyl, oxathiazinyl, e.g. 1,2,5- or 1,2,6-oxathiazinyl, 1,3,5-oxadiazinyl, dihydroisothiazolyl, dihydroisothiazole 1,1-dioxide, e.g. 2,3-dihydroisothiazole 1,1-dioxide, dihydropyrazinyl and tetrahydropyrazinyl groups.

The optional substituents which may be present on the cycloaliphatic, polycycloaliphatic, heterocycloaliphatic or heteropolycycloaliphatic groups represented by the group Cy¹ include one, two, three or more substituents selected from halogen atoms, and C₁₋₆ alkyl, e.g. methyl or ethyl, halo(C₁₋₆)alkyl, e.g. halomethyl or haloethyl such as

- 5 difluoromethyl or trifluoromethyl, optionally substituted by hydroxy, e.g. -C(OH)(CF₃)₂, C₁₋₆ alkoxy, e.g. methoxy or ethoxy, halo(C₁₋₆)alkoxy, e.g. halomethoxy or haloethoxy such as difluoromethoxy or trifluoromethoxy, thiol, C₁₋₆ alkylthio, e.g. methylthio or ethylthio, carbonyl (=O), thiocabonyl (=S), imino (=NR^{4a}) (where R^{4a} is an -OH group or a C₁₋₆ alkyl group) or -(Alk³)_vR⁵ groups, in which Alk³ is a straight or branched C₁₋₃
- 10 alkylene chain, v is zero or the integer 1, and R⁵ is a C₃₋₈ cycloalkyl, -OH, -SH, -N(R⁶)(R⁷) (in which R⁶ and R⁷ is each independently selected from a hydrogen atom and an optionally substituted alkyl or C₃₋₈ cycloalkyl group), -OR⁶, -SR⁶, -CN, -NO₂, -CO₂R⁶, -SOR⁶, -SO₂R⁶, -SO₃R⁶, -OCO₂R⁶, -C(O)R⁶, -OC(O)R⁶, -C(S)R⁶, -C(O)N(R⁶)(R⁷), -OC(O)N(R⁶)(R⁷), -N(R⁶)C(O)R⁷, -C(S)N(R⁶)(R⁷), -N(R⁶)C(S)R⁷, -SO₂N(R⁶)(R⁷),
- 15 -N(R⁶)SO₂R⁷, -N(R⁶)C(O)N(R⁷)(R⁸) (where R⁸ is as defined for R⁶), -N(R⁶)C(S)N(R⁷)(R⁸) or -N(R⁶)SO₂N(R⁷)(R⁸) group, or an optionally substituted aromatic or heteroaromatic group.

Particular examples of Alk³ chains include -CH₂-, -CH₂CH₂-, -CH₂CH₂CH₂- and -CH(CH₃)CH₂- chains.

- 20 When R⁵, R⁶, R⁷ and/or R⁸ is present as a C₃₋₈ cycloalkyl group it may be, for example, a cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl group. Optional substituents which may be present on such groups include, for example, one, two or three substituents, which may be the same or different, selected from halogen atoms, for example fluorine, chlorine, bromine or iodine atoms, and hydroxy or C₁₋₆ alkoxy groups,
- 25 e.g. methoxy, ethoxy or isopropoxy groups.

When the groups R⁶ and R⁷ or R⁷ and R⁸ are both alkyl groups these groups may be joined, together with the N atom to which they are attached, to form a heterocyclic ring. Such heterocyclic rings may be optionally interrupted by a further heteroatom or heteroatom-containing group selected from -O-, -S-, -N(R⁷)-, -C(O)- and -C(S)- groups.

- 30 Particular examples of such heterocyclic rings include piperidinyl, pyrazolidinyl, morpholinyl, thiomorpholinyl; pyrrolidinyl, imidazolidinyl and piperazinyl rings.

When R⁵ is an optionally substituted aromatic or heteroaromatic group it may be any such group as described hereinafter in relation to Cy¹.

Additionally, when the group Cy¹ is a heterocycloaliphatic or heteropolycycloaliphatic group containing one or more nitrogen atoms each nitrogen atom may be optionally substituted by a group -(L⁵)_p(Alk⁴)_qR⁹ in which L⁵ is a -C(O)-, -C(O)O-, -C(S)-, -S(O)₂-, -CON(R⁶)- or -SO₂N(R⁶)- group; p is zero or the integer 1; Alk⁴

5 is an optionally substituted aliphatic or heteroaliphatic chain; q is zero or the integer 1; and R⁹ is a hydrogen atom or an optionally substituted cycloaliphatic, heterocycloaliphatic, polycycloaliphatic, heteropolycycloaliphatic, aromatic or heteroaromatic group as herein described in relation to Cy¹.

When Alk⁴ is present as an aliphatic or heteroaliphatic chain it may be, for

10 example, any aliphatic or heteroaliphatic chain as hereinbefore described for Alk¹.

Optionally substituted aromatic groups represented by the group Cy¹ include, for example, monocyclic and bicyclic fused ring C₆₋₁₂ aromatic groups, such as phenyl, 1- or 2-naphthyl, 1- or 2-tetrahydronaphthyl, indanyl or indenyl groups.

Heteroaromatic groups represented by the group Cy¹ include, for example, C₁₋₉

15 heteroaromatic groups containing, for example, one, two, three or four heteroatoms selected from oxygen, sulphur and nitrogen atoms. In general, the heteroaromatic groups may be, for example, monocyclic or bicyclic fused ring heteroaromatic groups. Monocyclic heteroaromatic groups include, for example, five- or six-membered heteroaromatic groups containing one, two, three or four heteroatoms selected from

20 oxygen, sulphur and nitrogen atoms. Bicyclic heteroaromatic groups include, for example, eight- to thirteen-membered fused ring heteroaromatic groups containing one, two or more heteroatoms selected from oxygen, sulphur and nitrogen atoms.

Particular examples of heteroaromatic groups of these types include pyrrolyl, furyl, thienyl, imidazolyl, N-(C₁₋₆ alkyl)imidazolyl, oxazolyl, isoxazolyl, thiazolyl,

25 isothiazolyl, pyrazolyl, 1,2,3-triazolyl, 1,2,4-triazolyl, 1,2,3-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, 1,3,4-thiadiazolyl, pyridinyl, pyrimidinyl, pyridazinyl, pyrazinyl, 1,3,5-triazinyl, 1,2,4-triazinyl, 1,2,3-triazinyl, benzofuryl, (2,3-dihydro)benzofuryl, benzothienyl, (2,3-dihydro)benzothienyl, benzotriazolyl, indolyl, indolinyl, indazolinyl, benzimidazolyl, imidazo[1,2-a]pyridinyl, benzothiazolyl,

30 benzoxazolyl, benzisoxazolyl, benzopyranyl, (3,4-dihydro)benzopyranyl, quinazolinyl, quinoxalinyl, naphthyridinyl, imidazo[1,5-a]pyridinyl, imidazo[1,5-a]pyrazinyl, imidazo[1,5-c]pyrimidinyl, pyrido[3,4-b]pyridinyl, pyrido[3,2-b]pyridinyl, pyrido[4,3-b]pyridinyl, quinolinyl, isoquinolinyl, phthalazinyl, tetrazolyl, 5,6,7,8-

tetrahydroquinolinyl, 5,6,7,8-tetrahydroisoquinolinyl, imidyl, e.g. succinimidyl, phthalimidyl or naphthalimidyl such as 1,8-naphthalimidyl, pyrazolo[4,3-*d*]pyrimidinyl, furo[3,2-*d*]pyrimidinyl, thieno[3,2-*d*]pyrimidinyl, pyrrolo[3,2-*d*]pyrimidinyl, pyrazolo[3,2-*b*]pyridinyl, furo[3,2-*b*]pyridinyl, thieno[3,2-*b*]pyridinyl, pyrrolo[3,2-*b*]pyridinyl, thiazolo[3,2-*a*]pyridinyl, pyrido[1,2-*a*]pyrimidinyl, tetrahydroimidazo[1,2-*a*]pyrimidinyl and dihydroimidazo[1,2-*a*]pyrimidinyl groups.

Optional substituents which may be present on aromatic or heteroaromatic groups represented by the group Cy¹ include one, two, three or more substituents, each selected from an atom or group R¹⁰ in which R¹⁰ is R^{10a} or -L⁶Alk⁵(R^{10a})_r, where R^{10a} is a halogen atom, or an amino (-NH₂), substituted amino, nitro, cyano, hydroxy (-OH), substituted hydroxy, formyl, carboxy (-CO₂H), esterified carboxy, thiol (-SH), substituted thiol, -COR¹¹ (where R¹¹ is an -L⁶Alk³(R^{10a})_r, aryl or heteroaryl group), -CSR¹¹, -SO₃H, -SOR¹¹, -SO₂R¹¹, -SO₃R¹¹, -SO₂NH₂, -SO₂NHR¹¹, -SO₂N(R¹¹)₂, -CONH₂, -CSNH₂, -CONHR¹¹, -CSNHR¹¹, -CON(R¹¹)₂, -CSN(R¹¹)₂, -N(R¹²)SO₂R¹¹ (where R¹² is a hydrogen atom or a straight or branched alkyl group), -N(SO₂R¹¹)₂, -N(R¹²)SO₂NH₂, -N(R¹²)SO₂NHR¹¹, -N(R¹²)SO₂N(R¹¹)₂, -N(R¹²)COR¹¹, -N(R¹²)CONH₂, -N(R¹²)CONHR¹¹, -N(R¹²)CON(R¹¹)₂, -N(R¹²)CSNH₂, -N(R¹²)CSNHR¹¹, -N(R¹²)CSN(R¹¹)₂, -N(R¹²)CSR¹¹, -N(R¹²)C(O)OR¹¹, -SO₂NHet¹ (where -NHet¹ is an optionally substituted C₅₋₇ cyclic amino group optionally containing one or more other -O- or -S- atoms or -N(R¹²)-, -C(O)- or -C(S)- groups), -CONHet¹, -CSNHet¹, -N(R¹²)SO₂NHet¹, -N(R¹²)CONHet¹, -N(R¹²)CSNHet¹, -SO₂N(R¹²)Het² (where -Het² is an optionally substituted monocyclic C₅₋₇ carbocyclic group optionally containing one or more other -O- or -S- atoms or -N(R¹²)-, -C(O)-, -S(O)- or -S(O)₂- groups), -Het², -CON(R¹²)Het², -CSN(R¹²)Het², -N(R¹²)CON(R¹²)Het², -N(R¹²)CSN(R¹²)Het², -N(R¹²)SO₂N(R¹²)Het², aryl or heteroaryl group; L⁶ is a covalent bond or a linker atom or group as hereinbefore defined for L²; Alk⁵ is an optionally substituted straight or branched C₁₋₆ alkylene, C₂₋₆ alkenylene or C₂₋₆ alkynylene chain, optionally interrupted by one, two or three -O- or -S- atoms or -S(O)-, -S(O)₂- or -N(R¹²)-, e.g. -N(CH₃)-, groups; and r is zero or the integer 1, 2, or 3. It will be appreciated that when two R¹¹ or R¹² groups are present in one of the above substituents the R¹¹ and R¹² groups may be the same or different.

When in the group -L⁶Alk⁵(R^{10a})_r r is an integer 1, 2 or 3, it is to be understood that the substituent or substituents R^{10a} may be present on any suitable carbon atom in Alk⁵.

Where more than one R^{10a} substituent is present these may be the same or different and may be present on the same atom or on different atoms in Alk⁵. Clearly, when r is zero and no substituent R^{10a} is present the alkylene, alkenylene or alkynylene chain represented by Alk⁵ becomes an alkyl, alkenyl or alkynyl group.

5 When R^{10a} is a substituted amino group it may be, for example, a group -NHR¹¹ (where R¹¹ is as defined above) or a group -N(R¹¹)₂ wherein each R¹¹ group is the same or different.

When R^{10a} is a halogen atom it may be, for example, a fluorine, chlorine, bromine or iodine atom.

10 When R^{10a} is a substituted hydroxy or substituted thiol group it may be, for example, a group -OR¹¹ or -SR¹² respectively.

Esterified carboxy groups represented by the group R^{10a} include groups of formula -CO₂Alk⁶ wherein Alk⁶ is a straight or branched, optionally substituted C₁₋₈ alkyl group such as a methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl or *tert*-butyl group; a C₆₋₁₂ aryl(C₁₋₈ alkyl) group such as an optionally substituted benzyl, phenylethyl, phenylpropyl, 1-naphthylmethyl or 2-naphthylmethyl group; a C₆₋₁₂ aryl group such as an optionally substituted phenyl, 1-naphthyl or 2-naphthyl group; a C₆₋₁₂ aryloxy(C₁₋₈ alkyl) group such as an optionally substituted phenoxyethyl, phenoxyethyl, 1-naphthoxyethyl or 2-naphthoxyethyl group; an optionally substituted C₁₋₈ alkanoyloxy(C₁₋₈ alkyl) group, such as a pivaloyloxyethyl, propionyloxyethyl or propionyloxypropyl group; or a C₆₋₁₂ aroyloxy(C₁₋₈ alkyl) group such as an optionally substituted benzyloxyethyl or benzyloxypropyl group. Optional substituents present on the Alk⁶ group include R^{10a} atoms and groups as described above.

When Alk⁵ is present in or as a substituent it may be, for example, a -CH₂-,

25 -CH(CH₃)-, -C(CH₃)₂-, -CH₂CH₂-, -CH₂CH₂CH₂-, -CH(CH₃)CH₂-, -CH₂CH₂CH₂CH₂-,
 -CH₂CH(CH₃)CH₂-, -CH(CH₃)CH₂CH₂-, -C(CH₃)₂CH₂-, -CH=CH-, -CH=CHCH₂-,
 -CH₂CH=CH-, -CH=CHCH₂CH₂-, -CH₂CH=CHCH₂-, -CH₂CH₂CH=CH-, -C≡C-,
 -C≡CCH₂-, -CH₂C≡C-, -C≡CCH₂CH₂-, -CH₂C≡CCH₂- or -CH₂CH₂C≡C- chain, optionally interrupted by one, two or three -O- or -S- atoms or -S(O)-, -S(O)₂- or -N(R¹²)-, e.g.

30 -N(CH₃)-, groups. The aliphatic chains represented by Alk⁵ may be optionally substituted by one, two or three halogen atoms in addition to any R^{10a} groups that may be present.

Aryl or heteroaryl groups represented by the groups R^{10a} or R¹¹ include mono- and bicyclic optionally substituted C₆₋₁₂ aromatic or C₁₋₉ heteroaromatic groups as described

above for the group Cy¹. The aromatic and heteroaromatic groups may be attached to the group Cy¹ in compounds of formula (1) by any carbon atom or heteroatom, e.g. nitrogen atom, as appropriate.

It will be appreciated that when -NHet¹ or -Het² forms part of a substituent R¹⁰ the 5 heteroatoms or heteroatom-containing groups that may be present within the ring -NHet¹ or -Het² take the place of carbon atoms within the parent carbocyclic ring.

Thus, when -NHet¹ or -Het² forms part of a substituent R¹⁰ each may be, for example, an optionally substituted pyrrolidinyl, imidazolidinyl, pyrazolidinyl, piperazinyl, morpholinyl, thiomorpholinyl, piperidinyl or thiazolidinyl group. Additionally, -Het² may 10 represent, for example, an optionally substituted cyclopentyl or cyclohexyl group. Optional substituents which may be present on -NHet¹ include those substituents described above when Cy¹ is a heterocycloaliphatic group.

Particularly useful atoms or groups represented by R¹⁰ include fluorine, chlorine, bromine or iodine atoms, and C₁₋₆ alkyl, e.g. methyl, ethyl, n-propyl, isopropyl, n-butyl or 15 tert-butyl, optionally substituted phenyl, pyridinyl, pyrimidinyl, pyrrolyl, furyl, thiazolyl or thienyl, C₁₋₆ hydroxyalkyl, e.g. hydroxymethyl or hydroxyethyl, carboxy(C₁₋₆ alkyl), e.g. carboxyethyl, C₁₋₆ alkylthio, e.g. methylthio or ethylthio, carboxy(C₁₋₆ alkyl)thio, e.g. carboxymethylthio, 2-carboxyethylthio or 3-carboxypropylthio, C₁₋₆ alkoxy, e.g. methoxy or ethoxy, hydroxy(C₁₋₆ alkoxy), e.g. 2-hydroxyethoxy, optionally substituted phenoxy, 20 pyridinyloxy, thiazolyloxy, phenylthio or pyridinylthio, C₃₋₇ cycloalkyl, e.g. cyclobutyl or cyclopentyl, C₅₋₇ cycloalkoxy, e.g. cyclopentyloxy, halo(C₁₋₆ alkyl), e.g. trifluoromethyl, halo(C₁₋₆ alkoxy), e.g. trifluoromethoxy, C₁₋₆ alkylamino, e.g. methylamino, ethylamino, -CH(CH₃)NH₂ or -C(CH₃)₂NH₂, halo(C₁₋₆ alkyl)amino, e.g. fluoro(C₁₋₆ alkyl)amino such as -CH(CF₃)NH₂ or -C(CF₃)₂NH₂, amino (-NH₂), amino(C₁₋₆ alkyl), e.g. aminomethyl or 25 aminoethyl, di(C₁₋₆ alkyl)amino, e.g. dimethylamino or diethylamino, C₁₋₆ alkylamino(C₁₋₆ alkyl), e.g. ethylaminoethyl, di-(C₁₋₆ alkyl)amino(C₁₋₆ alkyl), e.g. diethylaminoethyl, amino(C₁₋₆ alkoxy), e.g. aminoethoxy, C₁₋₆ alkylamino(C₁₋₆ alkoxy), e.g. methylaminoethoxy, di(C₁₋₆ alkyl)amino(C₁₋₆ alkoxy), e.g. dimethylaminoethoxy, diethylaminoethoxy, diisopropylaminoethoxy or dimethylaminopropano, imido, such as phthalimido or 30 naphthalimido, e.g. 1,8-naphthalimido, nitro, cyano, hydroxy (-OH), formyl [HC(O)-], carboxy (-CO₂H), -CO₂Alk⁶ (where Alk⁶ is as defined above), C₁₋₆ alkanoyl, e.g. acetyl, optionally substituted benzoyl, thiol (-SH), thio(C₁₋₆ alkyl), e.g. thiomethyl or thioethyl, sulphonyl (-SO₃H), C₁₋₆ alkylsulphonyl, e.g. methylsulphonyl, aminosulphonyl (-SO₂NH₂),

C_{1-6} alkylaminosulphonyl, e.g. methylaminosulphonyl or ethylaminosulphonyl, di(C_{1-6} alkyl)aminosulphonyl, e.g. dimethylaminosulphonyl or diethylaminosulphonyl, phenylaminosulphonyl, carboxamido (-CONH₂), C_{1-6} alkylaminocarbonyl, e.g. methylaminocarbonyl or ethylaminocarbonyl, di(C_{1-6} alkyl)aminocarbonyl, e.g.

5 dimethylaminocarbonyl or diethylaminocarbonyl, amino(C_{1-6} alkyl)aminocarbonyl, e.g. aminoethylaminocarbonyl, di(C_{1-6} alkyl)amino(C_{1-6} alkyl)aminocarbonyl, e.g. diethylaminoethylaminocarbonyl, aminocarbonylamino, C_{1-6} alkylamino carbonylamino, e.g. methylamino carbonylamino or ethylamino carbonylamino, di(C_{1-6} alkyl)aminocarbonylamino, e.g. dimethylaminocarbonylamino or

10 diethylamino carbonylamino, C_{1-6} alkylaminocarbonyl(C_{1-6} alkyl)amino, e.g. methylaminocarbonylmethylamino, aminothiocarbonylamino, C_{1-6} alkylaminothiocarbonylamino, e.g. methylaminothiocarbonylamino or ethylaminothiocarbonylamino, di(C_{1-6} alkylaminothiocarbonylamino, e.g. dimethylaminothiocarbonylamino or diethylaminothiocarbonylamino, C_{1-6}

15 alkylaminothiocarbonyl(C_{1-6} alkyl)amino, e.g. ethylaminothiocarbonylmethylamino, -CONHC(=NH)NH₂, C_{1-6} alkylsulphonylamino, e.g. methylsulphonylamino or ethylsulphonylamino, di(C_{1-6} alkyl)sulphonylamino, e.g. dimethylsulphonylamino or diethylsulphonylamino, optionally substituted phenylsulphonylamino, aminosulphonylamino (-NHSO₂NH₂), C_{1-6} alkylaminosulphonylamino, e.g. methylaminosulphonylamino or

20 ethylaminosulphonylamino, di(C_{1-6} alkyl)aminosulphonylamino, e.g. dimethylamino-sulphonylamino or diethylaminosulphonylamino, optionally substituted morpholinylsulphonylamino or morpholinylsulphonyl(C_{1-6} alkyl)amino, optionally substituted phenylaminosulphonylamino, C_{1-6} alkanoylamino, e.g. acetylarnino, amino(C_{1-6} alkanoyl)arnino, e.g. aminoacetylarnino, di(C_{1-6} alkyl)arnino(C_{1-6} alkanoyl)arnino, e.g.

25 dimethylaminoacetylarnino, C_{1-6} alkanoylamino(C_{1-6} alkyl), e.g. acetylaminomethyl, C_{1-6} alkanoylamino(C_{1-6} alkyl)arnino, e.g. acetamidoethylarnino, C_{1-6} alkoxy carbonylamino, e.g. methoxycarbonylamino, ethoxycarbonylamino or *tert*-butoxycarbonylamino, or optionally substituted benzyloxy, pyridinylmethoxy, thiazolylmethoxy, benzyloxycarbonylamino, benzyloxycarbonylamino(C_{1-6} alkyl), e.g. benzyloxycarbonylaminoethyl, benzothio,

30 pyridinylmethylthio or thiazolylmethylthio groups.

A further particularly useful group of substituents represented by R¹⁰ when present on aromatic or heteroaromatic groups includes substituents of formula -L⁶Alk⁵R^{10a} where L⁶ is preferably a covalent bond, an -O- or -S- atom or a -N(R³)-, -C(O)-, -C(O)O-, -OC(O)-,

-N(R³)CO-, -CON(R³)- or -N(R³)S(O)₂- group; Alk⁵ is an optionally substituted C₁₋₆ alkylene group optionally interrupted by one or two -O- or -S- atoms or -N(R¹²)-, -C(O)-, -C(S)-, -CON(R¹²)- or -N(R¹²)CO- groups; and R^{10a} is an optionally substituted -Het² group as herein defined or an optionally substituted heteroaromatic group as hereinbefore described in
5 relation to Cy¹.

Where desired, two R¹⁰ substituents may be linked together to form a cyclic group such as a cyclic ether, e.g. a C₁₋₆ alkylene group such as methylenedioxy or ethylenedioxy.

It will be appreciated that where two or more R¹⁰ substituents are present, these
10 need not necessarily be the same atoms and/or groups. In general, the substituent(s) may be present at any available ring position on the aromatic or heteroaromatic group represented by the group Cy¹.

As used above, the terms "alkylamino" and "dialkylamino" are intended to include the groups -NHR¹³ and -N(R¹³)(R¹⁴) respectively, where R¹³ and R¹⁴ is each
15 independently an optionally substituted straight or branched alkyl group or both together with the N atom to which they are attached form an optionally substituted heterocycloalkyl group which may contain a further heteroatom or heteroatom-containing group such as an -O- or -S- atom or an -N(R^{2a})- group. Particular examples of such
20 optionally substituted heterocycloalkyl groups include optionally substituted pyrrolidinyl, piperidinyl, morpholinyl, thiomorpholinyl and N'-(C₁₋₆ alkyl)piperazinyl groups. The optional substituents which may be present on such heterocycloalkyl groups include those optional substituents as described hereinafter in relation to aliphatic chains such as Alk¹.

The optionally substituted aromatic or heteroaromatic group represented by Ar in compounds of the invention may be any aromatic or heteroaromatic group as hereinbefore
25 generally or particularly described for Cy¹. Optional substituents which may be present include those R¹⁰ atoms and groups as generally or particularly described in relation to Cy¹ aromatic and heteroaromatic groups.

Suitable values of R⁶ in the compounds of formula (1) above include hydrogen,
-CN, -COR¹, -CO₂R¹, -CONR^{1a}R^{2a}, -S(O)₂NR^{1a}R^{2a}, -CONR^{1a}OR^{2a} and
30 -C(O)N(R^{3a})NR^{1a}R^{2a}.

Suitably, R¹ represents methyl, ethyl or trifluoromethyl. In one embodiment, R¹ is ethyl. In another embodiment, R¹ is trifluoromethyl.

Suitable values of R^{1a} include hydrogen and methyl.

Suitable values of R^{2a} include hydrogen and methyl.

In a typical embodiment, R^{1a} and R^{2a} together represent $-(CH_2)_4-$. In another embodiment, R^{1a} and R^{2a} together represent $-CH(CH_2OH)(CH_2)_3-$.

Typically, R^{3a} is hydrogen.

5 Detailed values of R^e include hydrogen, cyano, trifluoroacetyl, ethoxycarbonyl, aminocarbonyl, methylaminocarbonyl, dimethylaminocarbonyl, pyrrolidinylcarbonyl, hydroxymethyl-pyrrolidinylcarbonyl, pyrrolidinylsulphonyl, (*N*-methoxy)(*N*-methyl)-aminocarbonyl and hydrazinocarbonyl.

In general, in compounds of formula (1) X and Y is each preferably a $-C(R^e)=$ or 10 $-C(Alk^2R^e)=$ group. R^e in these compounds is especially a hydrogen atom or a $-CONR^{1a}R^{2a}$ or $-S(O)_2NR^{1a}R^{2a}$ group. Alk^2 when present is in particular a $-CH_2-$ or $-(CH_2)_2-$ chain.

In one preference, X is a $-CH=$ group and Y is a $-C(R^e)=$ or $-C(Alk^2R^e)=$ group, especially where R^e is a hydrogen atom or a $-CONR^{1a}R^{2a}$ or $-S(O)_2NR^{1a}R^{2a}$ group and 15 Alk^2 is a $-CH_2-$ or $-(CH_2)_2-$ chain.

In one group of compounds of formula (1) the bond represented by the dashed line is present and A is a $-C(R^b)=$ group. In these compounds R^b is preferably a hydrogen atom or a C_{1-4} alkyl group, especially a methyl, ethyl, *n*-propyl or isopropyl group. More particularly, R^b is a methyl group or more especially a hydrogen atom.

20 One particularly useful class of compounds according to the invention has the formula (1) wherein A, X and Y is each a $-CH=$ group and R^a , Alk^1 , n , L^1 , Cy^1 , L and Ar are as generally or particularly defined herein; and the salts, solvates, hydrates and *N*-oxides thereof.

In general, in compounds of formula (1) R^a is preferably a hydrogen atom or a C_{1-4} 25 alkyl group, especially a methyl, ethyl, *n*-propyl or isopropyl group. In particular, R^a is a methyl group or more especially a hydrogen atom.

When in compounds of formula (1) n is the integer 1, Alk^1 is preferably an optionally substituted C_{1-6} alkylene chain, especially an optionally substituted $-CH_2-$, $-CH_2CH_2-$, $-CH_2CH_2CH_2-$, $-CH(CH_3)CH_2-$ or $-CH_2CH(CH_3)-$ chain, more especially a 30 $-CH_2-$ or $-CH_2CH_2-$ chain, and most especially a $-CH_2-$ chain.

In one class of compounds of formula (1) n is zero.

The group L^1 in compounds of formula (1) is preferably a covalent bond, an $-O-$ or $-S-$ atom or an $-N(R^3)-$, especially $-NH-$ or $-N(CH_3)-$, $-C(O)-$, $-C(S)-$, $-S(O)-$ or $-S(O)_2-$

group. More particularly, L¹ is a covalent bond, an -O- or -S- atom or a -NH- group. L¹ is most preferably a covalent bond.

Cy¹ in compounds of formula (1) is preferably an optionally substituted cycloaliphatic, aromatic or heteroaromatic group as hereinbefore generally and

5 particularly defined.

Particularly preferred Cy¹ optionally substituted cycloaliphatic groups include optionally substituted C₃₋₇ cycloalkyl groups, especially cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl groups.

Particularly preferred optional substituents which may be present on Cy¹

10 optionally substituted cycloaliphatic groups include halogen atoms, especially fluorine, chlorine or bromine atoms, C₁₋₆ alkyl groups, especially C₁₋₃ alkyl groups, most especially a methyl group, halo(C₁₋₆ alkyl) groups, especially fluoro(C₁₋₆ alkyl) groups, most especially a -CF₃ group, C₁₋₆ alkoxy groups, especially a methoxy, ethoxy, propoxy or isopropoxy group, and halo(C₁₋₆ alkoxy) groups, especially fluoro(C₁₋₆ alkoxy) groups,

15 most especially a -OCF₃ group, or a cyano (-CN), esterified carboxy, especially -CO₂CH₃ or -CO₂C(CH₃)₃, nitro (-NO₂), amino (-NH₂), substituted amino, especially -NHCH₃ or -N(CH₃)₂, -C(O)R⁶, especially -C(O)CH₃, or -N(R⁶)C(O)R⁷, especially -NHCOCH₃, group.

Particularly preferred Cy¹ aromatic groups include optionally substituted phenyl groups. Particularly preferred heteroaromatic groups include optionally substituted monocyclic heteroaromatic groups, especially optionally substituted five- or six-membered heteroaromatic groups containing one, two, three or four heteroatoms selected from oxygen, sulphur and nitrogen atoms. Particularly preferred optionally substituted monocyclic heteroaromatic groups include optionally substituted furyl, thienyl, pyrrolyl, oxazolyl, thiazolyl, pyridinyl, pyrimidinyl and triazinyl groups.

Particularly preferred optional substituents which may be present on Cy¹ aromatic or heteroaromatic groups include atoms or groups -R^{10a} and -L⁶Alk⁵(R^{10a}), as hereinbefore defined. Particularly useful optional substituents include halogen atoms, especially fluorine, chlorine or bromine atoms, C₁₋₆ alkyl groups, especially C₁₋₃ alkyl groups, most especially a methyl group, halo(C₁₋₆ alkyl) groups, especially fluoro(C₁₋₆ alkyl) groups, most especially a -CF₃ group, C₁₋₆ alkoxy groups, especially a methoxy, ethoxy, n-propoxy or isopropoxy group, and halo(C₁₋₆ alkoxy) groups, especially fluoro(C₁₋₆ alkoxy) groups, most especially a -OCF₃ group, or a cyano (-CN), carboxy (-CO₂H),

esterified carboxy ($-\text{CO}_2\text{Alk}^6$), especially $-\text{CO}_2\text{CH}_3$, $-\text{CO}_2\text{CH}_2\text{CH}_3$ or $-\text{CO}_2\text{C}(\text{CH}_3)_3$, nitro ($-\text{NO}_2$), amino ($-\text{NH}_2$), substituted amino, especially $-\text{NHCH}_3$ or $-\text{N}(\text{CH}_3)_2$, $-\text{COR}^{11}$, especially $-\text{COCH}_3$, or $-\text{N}(\text{R}^{12})\text{COR}^{11}$, especially $-\text{NHCOCH}_3$, group.

Further preferred optional substituents which may be present on Cy^1 aromatic or

- 5 heteroaromatic groups include groups of formula $-\text{L}^6\text{Alk}^5(\text{R}^{10a})_r$, in which r is the integer 1; L^6 is a covalent bond, an $-\text{O}-$ or $-\text{S}-$ atom, or a $-\text{N}(\text{R}^3)-$, especially $-\text{NH-}$ or $-\text{N}(\text{CH}_3)-$, $-\text{C}(\text{O})-$, $-\text{C}(\text{S})-$, $-\text{C}(\text{O})\text{O}-$, $-\text{OC}(\text{O})-$, $-\text{N}(\text{R}^3)\text{CO}-$, especially $-\text{NHCO-}$, or $-\text{CON}(\text{R}^3)-$, especially $-\text{CONH-}$, group; Alk^5 is a C_{1-6} alkylene chain, especially a $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{CH}_2-$ or $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2-$ chain; and R^{10a} is a substituted hydroxy group,
- 10 especially a $-\text{OCH}_3$, $-\text{OCH}_2\text{CH}_3$ or $-\text{OCH}(\text{CH}_3)_2$ group, a substituted amino group, especially a $-\text{N}(\text{CH}_3)_2$ or $-\text{N}(\text{CH}_2\text{CH}_3)_2$ group, or a $-\text{Het}^2$ group, especially an optionally substituted monocyclic C_{5-7} carbocyclic group containing one, two or three $-\text{O}-$, $-\text{S}-$, $-\text{N}(\text{R}^{12})-$, especially $-\text{NH-}$ or $-\text{N}(\text{CH}_3)-$, or $-\text{C}(\text{O})-$ groups within the ring structure as previously described, most especially an optionally substituted pyrrolidinyl,
- 15 imidazolidinyl, piperidinyl, e.g. *N*-methylpiperidinyl, morpholinyl, thiomorpholinyl or piperazinyl group, or R^{10a} is an optionally substituted heteroaromatic group, especially a five- or six-membered monocyclic heteroaromatic group containing one, two, three or four heteroatoms selected from oxygen, sulphur and nitrogen atoms, such as an optionally substituted pyrrolyl, furyl, thienyl, imidazolyl, triazolyl, pyridinyl, pyrimidinyl, triazinyl,
- 20 pyridazinyl or pyrazinyl group. Particularly preferred optional substituents on the $-\text{Het}^2$ groups just described include hydroxy ($-\text{OH}$) and carboxy ($-\text{CO}_2\text{H}$) groups or those preferred optional substituents just described in relation to the group Cy^1 .

In one particularly preferred group of compounds of formula (1) Cy^1 is an optionally substituted phenyl group, especially a phenyl group optionally substituted by

- 25 one, two or three optional substituents where at least one, and preferably two, optional substituents are located *ortho* to the bond joining Cy^1 to the remainder of the compound of formula (1). Particularly preferred *ortho* substituents include halogen atoms, especially fluorine or chlorine atoms, C_{1-3} alkyl groups, especially methyl, C_{1-3} alkoxy groups, especially methoxy, halo(C_{1-3} alkyl) groups, especially $-\text{CF}_3$, halo(C_{1-3} alkoxy) groups, especially $-\text{OCF}_3$, and cyano ($-\text{CN}$) groups. In this class of compounds a second or third optional substituent when present in a position other than the *ortho* positions of the ring Cy^1 may be preferably an atom or group $-\text{R}^{10a}$ or $-\text{L}^6\text{Alk}^5(\text{R}^{10a})_r$, as herein generally and particularly described.

Particular optional substituents on Cy¹ include C₁₋₆ alkyl and C₁₋₆ alkoxy. Specific optional substituents on Cy¹ include methyl and methoxy.

Specific values of Cy¹ include phenyl, methylphenyl (especially 4-methylphenyl), methoxyphenyl (especially 4-methoxyphenyl), thienyl (especially thien-3-yl) and indolyl

5 (especially indol-5-yl).

Particularly preferred Ar aromatic groups in compounds of formula (1) include optionally substituted phenyl groups. Particularly preferred heteroaromatic groups include optionally substituted monocyclic heteroaromatic groups, especially optionally substituted five- or six-membered heteroaromatic groups containing one, two, three or
10 four heteroatoms selected from oxygen, sulphur and nitrogen atoms. Particularly preferred optionally substituted monocyclic heteroaromatic groups include optionally substituted furyl, thienyl, pyrrolyl, oxazolyl, thiazolyl, pyridinyl, pyrimidinyl or triazinyl groups.

Particularly preferred optional substituents which may be present on Ar aromatic
15 or heteroaromatic groups include atoms or groups -R^{10a} or -L⁶Alk⁵(R^{10a}), as hereinbefore defined. Particularly useful optional substituents include halogen atoms, especially fluorine, chlorine or bromine atoms, C₁₋₆ alkyl groups, especially C₁₋₃ alkyl groups, most especially a methyl group, halo(C₁₋₆ alkyl) groups, especially fluoro(C₁₋₆ alkyl) groups, most especially a -CF₃ group, C₁₋₆ alkoxy groups, especially a methoxy, ethoxy, n-
20 propoxy or isopropoxy group, and halo(C₁₋₆ alkoxy) groups, especially fluoro(C₁₋₆ alkoxy) groups, most especially a -OCF₃ group, or a cyano (-CN), esterified carboxy, especially -CO₂CH₃ or -CO₂C(CH₃)₃, nitro (-NO₂), amino (-NH₂), substituted amino, especially -NHCH₃ or -N(CH₃)₂, -COR¹¹, especially -COCH₃, or -N(R¹²)COR¹¹, especially -NHCOCH₃, group.
25 Particularly useful Ar groups in compounds of formula (1) include phenyl and mono- or disubstituted phenyl groups in which each substituent is in particular a -R^{10a} or -L⁶Alk⁵(R^{10a}), atom or group as just defined and is especially a halogen atom or a C₁₋₃ alkyl, C₁₋₃ alkoxy or -CN group.

Suitable values of Ar include phenyl, pyridinyl, thienyl and benzothienyl, any of
30 which groups may be optionally substituted by one or more, typically by one or two, substituents. Examples of typical optional substituents which may be present on Ar include halogen (especially fluoro or chloro), cyano, C₁₋₆ alkyl (especially methyl), C₁₋₆ alkoxy (especially methoxy) and nitro.

Detailed values of Ar include phenyl, fluorophenyl, difluorophenyl, chlorophenyl, dichlorophenyl, (chloro)(fluoro)phenyl, cyanophenyl, methylphenyl, (fluoro)(methyl)-phenyl, methoxyphenyl, nitrophenyl, pyridinyl, chlorothienyl and benzothienyl.

Suitably, R^{1c} represents hydrogen or methyl, especially hydrogen. Suitably, R^{1f}

5 represents hydrogen.

In general, in compounds of formula (1) L is preferably a -CH₂- group.

In another embodiment, L is a -CH(CH₃)- group.

In a further embodiment, L is a -C(O)- group.

In an additional embodiment, L is a -CH₂CH₂- group.

10 Particularly useful compounds of the invention include each of the compounds described in the Examples hereinafter, and the salts, solvates, hydrates and N-oxides thereof.

Compounds according to the invention, including 3,7-dibenzyl-1,7-dihydro-2H-purinone, are potent and selective inhibitors of p38 kinases, including all isoforms and

15 splice variants thereof. More specifically, the compounds of the invention are inhibitors of p38 α , p38 β and p38 β 2. The ability of the compounds to act in this way may be simply determined by employing tests such as those described in the Examples hereinafter.

20 The compounds of formula (1) and 3,7-dibenzyl-1,7-dihydro-2H-purinone are of use in modulating the activity of p38 kinases and in particular are of use in the prophylaxis and treatment of any p38 kinase mediated diseases or disorders in a human or other mammal. The invention extends to such a use and to the use of the compounds for the manufacture of a medicament for treating such diseases or disorders. Furthermore, the invention extends to the administration to a human of an effective amount of a p38 inhibitor for treating any such disease or disorder.

25 The invention also extends to the prophylaxis or treatment of any disease or disorder in which p38 kinase plays a role including conditions caused by excessive or unregulated pro-inflammatory cytokine production, including for example excessive or unregulated TNF, IL-1, IL-6 and IL-8 production in a human or other mammal. The invention extends to such a use and to the use of the compounds for the manufacture of a medicament for treating such cytokine-mediated diseases or disorders. Furthermore, the invention extends to the administration to a human of an effective amount of a p38 inhibitor for treating any such disease or disorder.

Diseases or disorders in which p38 kinase plays a role either directly or via pro-inflammatory cytokines including the cytokines TNF, IL-1, IL-6 and IL-8 include without limitation autoimmune diseases, inflammatory diseases, destructive-bone disorders, proliferative disorders, neurodegenerative disorders, viral diseases, allergies, infectious diseases, heart attacks, angiogenic disorders, reperfusion/ischemia in stroke, vascular hyperplasia, organ hypoxia, cardiac hypertrophy, thrombin-induced platelet aggregation and conditions associated with prostaglandin endoperoxidase synthetase-2 (COX-2).

5 Autoimmune diseases which may be prevented or treated include but are not limited to rheumatoid arthritis, inflammatory bowel disease, ulcerative colitis, Crohn's disease, multiple sclerosis, diabetes, glomerulonephritis, systemic lupus erythematosus, scleroderma, chronic thyroiditis, Grave's disease, hemolytic anemia, autoimmune gastritis, autoimmune neutropenia, thrombocytopenia, chronic active hepatitis, myasthenia gravis, atopic dermatitis, graft vs host disease and psoriasis.

10 The invention further extends to the particular autoimmune disease rheumatoid arthritis.

Inflammatory diseases which may be prevented or treated include but are not limited to asthma, allergies, respiratory distress syndrome, and acute or chronic pancreatitis.

15 Destructive bone disorders which may be prevented or treated include but are not limited to osteoporosis, osteoarthritis and multiple myeloma-related bone disorder.

Proliferative diseases which may be prevented or treated include but are not limited to acute or chronic myelogenous leukemia, Kaposi's sarcoma, metastatic melanoma and multiple myeloma.

20 Neurodegenerative diseases which may be prevented or treated include but are not limited to Parkinson's disease, Alzheimer's disease, cerebral ischemias and neurodegenerative disease caused by traumatic injury.

Viral diseases which may be prevented or treated include but are not limited to acute hepatitis infection (including hepatitis A, hepatitis B and hepatitis C), HIV infection and CMV retinitis.

25 Infectious diseases which may be prevented or treated include but are not limited to septic shock, sepsis and Shigellosis.

In addition, p38 inhibitors of this invention exhibit inhibition of expression of inducible pro-inflammatory proteins such as prostaglandin endoperoxidase synthetase-2,

otherwise known as cyclooxygenase-2 (COX-2), and are therefore of use in therapy. Pro-inflammatory mediators of the cyclooxygenase pathway derived from arachidonic acid are produced by inducible COX-2 enzyme. Regulation of COX-2 would regulate these pro-inflammatory mediators such as prostaglandins, which affect a wide variety of cells

5 and are important and critical inflammatory mediators of a wide variety of disease states and conditions. In particular, these inflammatory mediators have been implicated in pain, such as in the sensitization of pain receptors, or edema. Accordingly, additional p38-mediated conditions which may be prevented or treated include edema, analgesia, fever and pain such as neuromuscular pain, headache, dental pain, arthritis pain and pain caused

10 by cancer.

More particularly, as a result of their p38 inhibitory activity, compounds of the invention have utility in the prevention and treatment of diseases associated with cytokine production including but not limited to those diseases associated with TNF, IL-1, IL-6 and IL-8 production.

15 Thus, TNF-mediated diseases or conditions include for example rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, gouty arthritis and other arthritic conditions, sepsis, septic shock syndrome, adult respiratory distress syndrome, cerebral malaria, chronic pulmonary inflammatory disease, silicosis, pulmonary sarcoidosis, bone resportion disease, reperfusion injury, graft vs host reaction, allograft rejections, fever and

20 myalgias due to infection, cachexia secondary to infection, AIDS, ARC or malignancy, keloid formation, scar tissue formation, Crohn's disease, ulcerative colitis, pyresis, and viral infections such as HIV, CMV, influenza and herpes; veterinary viral infections such as lentivirus infections, including but not limited to equine infectious anemia virus, caprine arthritis virus, visna virus or maedi virus; and retrovirus infections, including

25 feline immunodeficiency virus, bovine immunodeficiency virus and canine immunodeficiency virus.

Compounds of the invention may also be used in the treatment of viral infections, where such viruses elicit TNF production *in vivo* or are sensitive to upregulation by TNF. Such viruses include those that produce TNF as a result of infection and those that are

30 sensitive to inhibition, for instance as a result of decreased replication, directly or indirectly by the TNF-inhibiting compounds of the invention. Such viruses include, but are not limited to, HIV-1, HIV-2 and HIV-3, Cytomegalovirus (CMV), influenza, adenovirus and the herpes group of viruses such as *Herpes zoster* and *Herpes simplex*.

IL-1 mediated diseases or conditions include for example rheumatoid arthritis, osteoarthritis, psoriatic arthritis, traumatic arthritis, rubella arthritis, inflammatory bowel disease, stroke, endotoxemia and/or toxic shock syndrome, inflammatory reaction induced by endotoxin, diabetes, pancreatic β -cell disease, Alzheimer's disease,

5 tuberculosis, atherosclerosis, muscle degeneration and cachexia.

IL-8 mediated diseases and conditions include for example those characterized by massive neutrophil infiltration such as psoriasis, inflammatory bowel disease, asthma, cardiac, brain and renal reperfusion injury, adult respiratory distress syndrome, thrombosis and glomerulonephritis. The increased IL-8 production associated with each
10 of these diseases is responsible for the chemotaxis of neutrophils into inflammatory sites. This is due to the unique property of IL-8 (in comparison to TNF, IL-1 and IL-6) of promoting neutrophil chemotaxis and activation. Therefore, inhibition of IL-8 production would lead to a direct reduction in neutrophil infiltration.

It is also known that both IL-6 and IL-8 are produced during rhinovirus (HRV)
15 infections and contribute to the pathogenesis of the common cold and exacerbation of asthma associated with HRV infection [Turner *et al.*, *Clin. Infect. Dis.*, 1997, **26**, 840; Grunberg *et al.*, *Am. J. Crit. Care Med.*, 1997, **155**, 1362; Zhu *et al.*, *J. Clin. Invest.*, 1996, **97**, 421]. It has also been demonstrated *in vitro* that infection of pulmonary epithelial cells (which represent the primary site of infection by HRV) with HRV results
20 in production of IL-6 and IL-8 [Sabauste *et al.*, *J. Clin. Invest.*, 1995, **96**, 549]. Therefore, p38 inhibitors of the invention may be used for the treatment or prophylaxis of the common cold or respiratory viral infection caused by human rhinovirus infection (HRV), other enteroviruses, coronavirus, influenza virus, parainfluenza virus, respiratory syncytial virus or adenovirus.

25 For the prophylaxis or treatment of a p38 or pro-inflammatory cytokine mediated disease the compounds according to the invention may be administered to a human or mammal as pharmaceutical compositions, and according to a further aspect of the invention we provide a pharmaceutical composition which comprises a compound of formula (1) together with one or more pharmaceutically acceptable carriers, excipients or
30 diluents.

Pharmaceutical compositions according to the invention may take a form suitable for oral, buccal, parenteral, nasal, topical, ophthalmic or rectal administration, or a form suitable for administration by inhalation or insufflation.

For oral administration, the pharmaceutical compositions may take the form of, for example, tablets, lozenges or capsules prepared by conventional means with pharmaceutically acceptable excipients such as binding agents (e.g. pregelatinised maize starch, polyvinylpyrrolidone or hydroxypropyl methyl cellulose); fillers (e.g. lactose,

5 microcrystalline cellulose or calcium hydrogenphosphate); lubricants (e.g. magnesium stearate, talc or silica); disintegrants (e.g. potato starch or sodium glycollate); or wetting agents (e.g. sodium lauryl sulphate). The tablets may be coated by methods well known in the art. Liquid preparations for oral administration may take the form of, for example, solutions, syrups or suspensions, or they may be presented as a dry product for

10 constitution with water or other suitable vehicle before use. Such liquid préparations may be prepared by conventional means with pharmaceutically acceptable additives such as suspending agents, emulsifying agents, non-aqueous vehicles or preservatives. The preparations may also contain buffer salts, flavouring agents, colouring agents or sweetening agents, as appropriate.

15 Preparations for oral administration may be suitably formulated to give controlled release of the active compound.

For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

The compounds of formula (1) may be formulated for parenteral administration by

20 injection, e.g. by bolus injection or infusion. Formulations for injection may be presented in unit dosage form, e.g. in glass ampoules or multi-dose containers, e.g. glass vials. The compositions for injection may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilising, preserving and/or dispersing agents. Alternatively, the active ingredient may

25 be in powder form for constitution with a suitable vehicle, e.g. sterile pyrogen-free water, before use.

In addition to the formulations described above, the compounds of formula (1) may also be formulated as a depot preparation. Such long-acting formulations may be administered by implantation or by intramuscular injection.

30 For nasal administration or administration by inhalation, the compounds according to the present invention may be conveniently delivered in the form of an aerosol spray presentation for pressurised packs or a nebuliser, with the use of a suitable propellant, e.g.

dichlorodifluoromethane, fluorotrichloromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas or mixture of gases.

The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The 5 pack or dispensing device may be accompanied by instructions for administration.

For topical administration the compounds according to the present invention may be conveniently formulated in a suitable ointment containing the active component suspended or dissolved in one or more pharmaceutically acceptable carriers. Particular carriers include, for example, mineral oil, liquid petroleum, propylene glycol, 10 polyoxyethylene, polyoxypropylene, emulsifying wax and water. Alternatively, the compounds according to the present invention may be formulated in a suitable lotion containing the active component suspended or dissolved in one or more pharmaceutically acceptable carriers. Particular carriers include, for example, mineral oil, sorbitan monostearate, polysorbate 60, cetyl esters wax, cetearyl alcohol, benzyl alcohol, 2-octyldodecanol and water. 15

For ophthalmic administration the compounds according to the present invention may be conveniently formulated as microionized suspensions in isotonic, pH-adjusted sterile saline, either with or without a preservative such as a bactericidal or fungicidal agent, for example phenylmercuric nitrate, benzylalkonium chloride or chlorhexidine 20 acetate. Alternatively, for ophthalmic administration compounds may be formulated in an ointment such as petrolatum.

For rectal administration the compounds according to the present invention may be conveniently formulated as suppositories. These can be prepared by mixing the active component with a suitable non-irritating excipient which is solid at room temperature but 25 liquid at rectal temperature and so will melt in the rectum to release the active component. Such materials include, for example, cocoa butter, beeswax and polyethylene glycols.

The quantity of a compound of the invention required for the prophylaxis or treatment of a particular condition will vary depending on the compound chosen and the condition of the patient to be treated. In general, however, daily dosages may range from 30 around 10 ng/kg to 1000 mg/kg, typically from 100 ng/kg to 100 mg/kg, e.g. around 0.01 mg/kg to 40 mg/kg body weight for oral or buccal administration, from around 10 ng/kg to 50 mg/kg body weight for parenteral administration, and from around 0.05 mg to

around 1000 mg, e.g. from around 0.5 mg to around 1000 mg, for nasal administration or administration by inhalation or insufflation.

The compounds of the invention may be prepared by a number of processes as generally described below and more specifically in the Examples hereinafter. In the

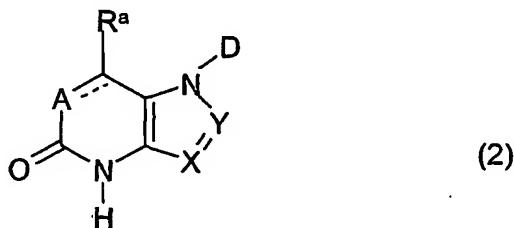
5 following process description, the symbols Ar, Cy¹, Alk¹, n, L¹, L, R^a, A, X and Y when used in the formulae depicted are to be understood to represent those groups described above in relation to formula (1) unless otherwise indicated. In the reactions described below, it may be necessary to protect reactive functional groups, for example hydroxy, amino, thio or carboxy groups, where these are desired in the final product, to avoid their

10 unwanted participation in the reactions. Conventional protecting groups may be used in accordance with standard practice [see, for example, Greene, T.W. and Wuts, P.G.M., *Protective Groups in Organic Synthesis*, John Wiley and Sons, 3rd edition, 1999]. In some instances, deprotection may be the final step in the synthesis of a compound of formula (1) and the processes according to the invention described hereinafter are to be

15 understood to extend to such removal of protecting groups.

Thus, according to a further aspect of the invention, a compound of formula (1) in which Cy¹ is as previously defined, but is other than an aromatic or heteroaromatic group when n is zero and L¹ is a covalent bond, may be prepared by alkylation of a compound of formula (2):

20



(where D is a hydrogen atom, a group -LAr, or a protecting group), with an alkylating agent of formula Cy¹L¹(Alk¹)_nZ, where Cy¹ is as just defined and Z is a leaving group

25 such as a halogen atom, e.g. a chlorine, bromine or iodine atom, or a sulphonyloxy group such as an alkylsulphonyloxy, e.g. trifluoromethylsulphonyloxy, or arylsulphonyloxy, e.g. phenylsulphonyloxy, group.

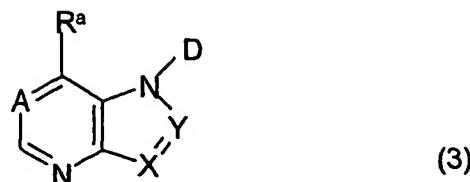
The reaction may be performed in the presence of a solvent, for example a substituted amide such as N,N-dimethylformamide, a cyclic ether such as tetrahydrofuran,

a sulphoxide such as dimethylsulphoxide, or mixtures thereof, optionally in the presence of a base, for example an inorganic base such as sodium hydride or potassium carbonate, or an organic base such as an organic amine, e.g. a cyclic amine such as 1,5-diazabicyclo[4.3.0]non-5-ene, or a resin-bound organic amine such as resin-bound 2-*tert*-butylimino-2-diethylamino-1,3-dimethylperhydro-1,3,2-diazaphosphorine (PS-BEMP), if necessary at an elevated temperature, for example 60 to 100°C.

Intermediates of formula (2) may also be used in another process according to the invention to prepare a compound of formula (1) in which n is zero, L¹ is a covalent bond and Cy¹ is an optionally substituted aromatic or heteroaromatic group by the reaction of 10 the intermediate with a boronic acid of formula Cy¹B(OH)₂ in which Cy¹ is as just defined. The reaction may be performed in an organic solvent, for example a halogenated hydrocarbon such as dichloromethane or dichloroethane, in the presence of a copper reagent, for example a copper(II) reagent such as copper(II) acetate, optionally in the presence of an oxidant, for example 2,2,6,6-tetramethyl-1-piperidinyloxy or pyridine N-oxide, optionally in the presence of a base, for example an organic amine such as an alkylamine, e.g. triethylamine, or an aromatic amine, e.g. pyridine, at a temperature from around ambient to the reflux temperature [see for example Chan, D.T. et al., *Tetrahedron Lett.*, 1998, 2933; Lam, P.Y.S. et al., *Tetrahedron Lett.*, 2001, 3415].

Moreover, for the preparation of a compound of formula (1) in which n is zero, L¹ 20 is a covalent bond and Cy¹ is a phenyl moiety substituted in the *ortho* and/or *para* positions by one, two or three nitro groups, the intermediate of formula (2) may be reacted with the appropriate fluoro-substituted nitrobenzene derivative. For example, where Cy¹ is 2-nitrophenyl, intermediate (2) will be reacted with 2-fluoronitrobenzene. The reaction is conveniently effected at an elevated temperature in the presence of a 25 strong base, e.g. sodium hydride, typically in a dipolar aprotic solvent such as *N,N*-dimethylformamide.

Intermediates of formula (2) are either known compounds (see, for example, Andresen et al., *ibid.*) or may be prepared in a multi-step process, firstly by oxidation of the corresponding pyridines or pyrimidines of formula (3):



with an oxidising agent such as hydrogen peroxide in the presence of an acid such as acetic acid, at an elevated temperature, for example around 70°C to 80°C, or alternatively

5 by reaction with a peracid such as peracetic acid or *m*-chloroperoxybenzoic acid in a solvent such as an ester, e.g. ethyl acetate, a halogenated hydrocarbon, e.g. dichloromethane, or an alcohol, e.g. *tert*-butanol, at a temperature from the ambient temperature to the reflux temperature, to yield the corresponding *N*-oxide; followed by sequential reaction with an anhydride, for example acetic anhydride, at an elevated

10 temperature, such as the reflux temperature, and then with an inorganic base, for example a carbonate such as aqueous potassium carbonate, in a solvent such as an ether, for example a cyclic ether, e.g. tetrahydrofuran, at around ambient temperature, to yield the desired intermediate of formula (2). Alternatively the *N*-oxide may be treated with trifluoroacetic anhydride in *N,N*-dimethylformamide from 0°C to ambient temperature to

15 yield the desired intermediate [see, for example, Kommo *et al.*, *Heterocycles*, 1986, 24, 2169].

Intermediates of formula (3) in which D is a protecting group can be advantageously used in the above reactions to generate a variety of compounds of the invention. Thus, for example, an intermediate of formula (3) in which D is a phenylsulphonyl group may be oxidised, converted to the corresponding pyridinone or pyrimidinone of formula (2), and then alkylated or arylated according to the reactions described above. At this point the phenylsulphonyl group can be removed, for example by treatment with a base such as sodium hydroxide in an alcohol such as methanol at ambient temperature. The resulting deprotected intermediate can then be reacted, in another process according to the invention, with a compound ArLZ (where Z is a leaving group as defined above) in the presence of a base and suitable solvent, as described above for the alkylation of intermediates of formula (2), to yield a desired compound of formula (1).

Where in the general processes described above intermediates of formula (3),

30 other intermediates such as alkylating agents of formula $Cy^1 L^1 (Alk^1)_n Z$, and any other

intermediates required in the synthesis of compounds of the invention, are not available commercially or known in the literature, they may be readily obtained from simpler known compounds by one or more standard synthetic methods employing substitution, oxidation, reduction or cleavage reactions. Particular substitution approaches include

5 conventional alkylation, arylation, heteroarylation, acylation, thioacetylation, halogenation, sulphonylation, nitration, formylation and coupling procedures. It will be appreciated that these methods may also be used to obtain or modify other intermediates and in particular compounds of formula (1) where appropriate functional groups exist in these compounds.

Thus, for example, intermediates of formula (3) where D is a protecting group

10 such as an arylsulphonyl group may be obtained from the corresponding $1H$ -compounds by reaction with a compound ArSO_2Hal (where Hal is a halogen atom such as a chlorine atom) in the presence of a base such as sodium hydride in a solvent such as dimethylsulphoxide at ambient temperature. Similarly, compounds of the invention and intermediates thereto generally may be obtained by sulphonylation of a compound

15 containing an -OH group by reaction with one of the above alkylating agents but in which Z is replaced by a -S(O)Hal or -SO₂Hal group in the presence of a base, for example an inorganic base such as sodium hydride, in a solvent such as an amide, e.g. a substituted amide such as *N,N*-dimethylformamide, at, for example, ambient temperature.

Compounds of the invention and intermediates thereto where A represents a

20 -N(R^b)- or -C(R^b)(R^c)- group may be generated from corresponding compounds of the invention or intermediates thereto where A represents a -N= or -C(R^b)= group by reduction, for instance by catalytic hydrogenation using a metal catalyst such as palladium on charcoal in the presence of hydrogen gas at an elevated pressure in a solvent such as an alcohol, e.g. ethanol, optionally at an elevated temperature, e.g. between 40°C

25 and 60°C.

Aromatic halogen substituents in the compounds of the invention may be subjected to halogen-metal exchange with a base, for example a lithium base such as *n*-butyllithium or *tert*-butyllithium, optionally at a low temperature, e.g. around -78°C, in a solvent such as tetrahydrofuran, and then quenched with an electrophile to introduce a

30 desired substituent. Thus, for example, a formyl group may be introduced by using *N,N*-dimethylformamide as the electrophile, a thiomethyl group may be introduced by using dimethyldisulphide as the electrophile, an alcohol group may be introduced by using an aldehyde as the electrophile, and an acid may be introduced by using carbon dioxide as

the electrophile. Aromatic acids of formula ArCO₂H may also be generated by quenching Grignard reagents of formula ArMgHal with carbon dioxide.

Aromatic acids of formula ArCO₂H generated by this method and acid-containing compounds in general may be converted to activated derivatives, e.g. acid halides, by

- 5 reaction with a halogenating agent such as a thionyl halide, e.g. thionyl chloride, a phosphorus trihalide such as phosphorus trichloride, or a phosphorus pentahalide such as phosphorus pentachloride, optionally in an inert solvent such as an aromatic hydrocarbon, e.g. toluene, or a chlorinated hydrocarbon, e.g. dichloromethane, at a temperature from about 0°C to the reflux temperature, or may be converted into Weinreb amides of formula
- 10 ArC(O)N(OMe)Me by direct treatment with the amine of formula HN(OMe)Me and a condensing agent such as EDC (*vide infra*), typically in the presence of a base such as triethylamine in a solvent such as dichloromethane; or by conversion to the acid halide as just described and subsequent reaction with the amine of formula HN(OMe)Me or a salt thereof, optionally in the presence of a base such as an organic amine, e.g. triethylamine,
- 15 in an inert solvent such as an aromatic hydrocarbon, e.g. toluene, or a chlorinated hydrocarbon, e.g. dichloromethane, at a temperature from about 0°C to ambient temperature.

Acid (-CO₂H) groups in the compounds of the invention or intermediates thereto may be transformed into amide (-CONH₂) groups by treatment with ammonia in the presence of a condensing agent, for example 1,1'-carbonyldiimidazole, typically in a solvent such as *N,N*-dimethylformamide. Similarly, acid (-CO₂H) groups in the compounds of the invention or intermediates thereto may be transformed into hydrazide (-CONHNH₂) groups by treatment with hydrazine hydrate, suitably in an alcoholic solvent such as 2-ethoxyethanol, typically at an elevated temperature.

- 25 Amide (-CONH₂) groups in the compounds of the invention or intermediates thereto may be transformed into cyano (-CN) groups by treatment with trifluoroacetic anhydride in the presence of pyridine and a solvent, e.g. a chlorinated solvent such as dichloromethane.

Compounds of formula (1) wherein R^e represents -CO₂H may be converted into

- 30 the corresponding compound wherein R^e represents -CONR^{1a}R^{2a} by conversion into an activated derivative, e.g. an acid halide, as described above, typically in a dipolar aprotic solvent such as *N,N*-dimethylformamide; followed by treatment with the appropriate amine of formula H-NR^{1a}R^{2a}, typically in an inert solvent such as tetrahydrofuran.

Alternatively, a compound of formula (1) wherein R^e represents -CO₂H may be converted directly into the corresponding compound wherein R^e represents -CONR^{1a}R^{2a} by treatment with the appropriate amine of formula H-NR^{1a}R^{2a} and a condensing agent such as EDC (*vide infra*), suitably in the presence of a catalyst such as 1-hydroxybenzotriazole,

5 typically in an inert solvent, e.g. a chlorinated solvent such as dichloromethane.

Compounds of the invention and intermediates thereto may be prepared by alkylation, arylation or heteroarylation. For example, compounds containing a -L¹H group (where L¹ is a linker atom or group) may be treated with an alkylating agent Cy¹Z² in which Z² is a leaving atom or group such as a halogen atom, e.g. a fluorine, chlorine, 10 bromine or iodine atom, or a sulphonyloxy group such as an alkylsulphonyloxy, e.g. trifluoromethylsulphonyloxy, or arylsulphonyloxy, e.g. *p*-toluenesulphonyloxy, group.

The reaction may be carried out in the presence of a base such as a carbonate, e.g. caesium carbonate or potassium carbonate, an alkoxide, e.g. potassium *tert*-butoxide, or a hydride, e.g. sodium hydride, in a dipolar aprotic solvent such as an amide, e.g. a 15 substituted amide such as *N,N*-dimethylformamide, or an ether, e.g. a cyclic ether such as tetrahydrofuran.

In another example, compounds containing a -L²H group as defined above may be functionalised by acylation or thioacetylation, for example by reaction with the alkylating agents just described but in which Z² is replaced by a -C(O)Z³, -C(S)Z³, -N(R²)C(O)Z³ or 20 -N(R²)C(S)Z³ group in which Z³ is a leaving atom or group as described for Z². The reaction may be performed in the presence of a base, such as a hydride, e.g. sodium hydride, or an amine, e.g. triethylamine or *N*-methylmorpholine, in a solvent such as a halogenated hydrocarbon, e.g. dichloromethane or carbon tetrachloride, or an amide, e.g. *N,N*-dimethylformamide, at for example ambient temperature. Alternatively, the 25 acylation may be carried out under the same conditions with an acid (for example one of the alkylating agents described above in which Z² is replaced by a -CO₂H group) in the presence of a condensing agent, for example a diimide such as 1-(3-dimethylamino-propyl)-3-ethylcarbodiimide or *N,N*'-dicyclohexylcarbodiimide, or a benzotriazole such as O-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate, 30 advantageously in the presence of a catalyst such as a *N*-hydroxy compound, e.g. a *N*-hydroxytriazole such as 1-hydroxybenzotriazole. Alternatively, the acid may be reacted with a chloroformate, for example ethyl chloroformate, prior to the desired acylation reaction.

In another example, compounds containing a $-L^2H$ group as defined above may be coupled with one of the alkylation agents just described but in which Z^2 is replaced by an $-OH$ group in a solvent such as tetrahydrofuran in the presence of a phosphine, e.g. triphenylphosphine, and an activator such as diethyl, diisopropyl or dimethyl

5 azodicarboxylate.

Ester groups such as $-CO_2Alk^6$, $-CO_2R^4$ and $-CO_2R^1$ (where R^1 is other than hydrogen) in the compounds of formula (1) and intermediates thereto may be converted to the corresponding acid ($-CO_2H$) by acid- or base-catalysed hydrolysis depending on the nature of the group Alk^6 , R^4 or R^1 . Acid- or base-catalysed hydrolysis may be achieved,

10 for example, by treatment with an organic or inorganic acid, e.g. trifluoroacetic acid, in an organic solvent, e.g. dichloromethane, or a mineral acid such as hydrochloric acid in a solvent such as 1,4-dioxane, or an alkali metal hydroxide, e.g. lithium hydroxide or sodium hydroxide, in an aqueous alcohol, e.g. aqueous methanol or aqueous ethanol.

In a further example, $-OR^6$ (where R^6 represents an alkyl group such as methyl) in compounds of formula (1) and intermediates thereto may be cleaved to the corresponding alcohol $-OH$ by reaction with boron tribromide in a solvent such as a halogenated hydrocarbon, e.g. dichloromethane, at a low temperature, e.g. around $-78^\circ C$.

Alcohol ($-OH$) groups may also be obtained by hydrogenation of a corresponding $-OCH_2R^{31}$ group (where R^{31} is an aryl group) using a metal catalyst, for example palladium on a support such as carbon, in a solvent such as ethanol in the presence of ammonium formate, cyclohexadiene or hydrogen, from around ambient temperature to the reflux temperature. In another example, $-OH$ groups may be generated from the corresponding ester (e.g. $-CO_2Alk^6$) or aldehyde ($-CHO$) by reduction, using for example a complex metal hydride such as lithium aluminium hydride or sodium borohydride in a solvent such as methanol.

In another example, $-OH$ groups in the compounds may be converted to a corresponding $-OR^6$ group by coupling with a reagent R^6OH in a solvent such as tetrahydrofuran in the presence of a phosphine, e.g. triphenylphosphine and an activator such as diethyl, diisopropyl or dimethyl azodicarboxylate.

30 Aminosulphonylamino ($-NHSO_2NH_2$) groups in the compounds may be obtained, in another example, by reaction of a corresponding amine ($-NH_2$) with sulphamide in the presence of an organic base such as pyridine at an elevated temperature, e.g. the reflux temperature.

In another example, compounds containing a $-NHCSR^7$ or $-CSNHR^7$ group may be prepared by treating a corresponding compound containing a $-NHCOR^7$ or $-CONHR^7$ group with a thiation reagent, such as Lawesson's Reagent or P_2S_5 , in an anhydrous solvent, for example a cyclic ether such as tetrahydrofuran, at an elevated temperature
5 such as the reflux temperature.

In a further example, amine ($-NH_2$) groups may be alkylated using a reductive alkylation process employing an aldehyde and a reducing agent. Suitable reducing agents include borohydrides, for example sodium triacetoxyborohydride or sodium cyanoborohydride. The reduction may be carried out in a solvent such as a halogenated
10 hydrocarbon, e.g. dichloromethane, a ketone such as acetone, or an alcohol, e.g. ethanol, where necessary in the presence of an acid such as acetic acid at around ambient temperature. Alternatively, the amine and aldehyde may be initially reacted in a solvent such as an aromatic hydrocarbon, e.g. toluene, and then subjected to hydrogenation in the presence of a metal catalyst, for example palladium on a support such as carbon, in a
15 solvent such as an alcohol, e.g. ethanol.

Amine ($-NH_2$) groups in the compounds of the invention or intermediates thereto may generally be transformed into halogen atoms, e.g. bromo, by treatment with a nitrite reagent, e.g. *tert*-butyl nitrite, in the presence of a copper(II) halide, e.g. copper(II) bromide, typically in a solvent such as acetonitrile.

20 In a further example, amine ($-NH_2$) groups in compounds of formula (1) and intermediates thereto may be obtained by hydrolysis from a corresponding imide by reaction with hydrazine in a solvent such as an alcohol, e.g. ethanol, at ambient temperature.

25 In another example, a nitro ($-NO_2$) group may be reduced to an amine ($-NH_2$), for example by catalytic hydrogenation using for example hydrogen in the presence of a metal catalyst, for example palladium on a support such as carbon, in a solvent such as an ether, e.g. tetrahydrofuran, or an alcohol, e.g. methanol, or by chemical reduction using for example a metal, e.g. tin or iron, in the presence of an acid such as hydrochloric acid.

30 In a further example, amine ($-CH_2NH_2$) groups in compounds of formula (1) and intermediates thereto may be obtained by reduction of nitriles ($-CN$), for example by catalytic hydrogenation using for example hydrogen in the presence of a metal catalyst, for example palladium on a support such as carbon, or Raney[®] nickel, in a solvent such as an ether, e.g. a cyclic ether such as tetrahydrofuran, or an alcohol, e.g. methanol or

ethanol, optionally in the presence of ammonia solution at a temperature from ambient temperature to the reflux temperature, or by chemical reduction using for example a metal hydride, e.g. lithium aluminium hydride, in a solvent such as an ether, e.g. a cyclic ether such as tetrahydrofuran, at a temperature from 0°C to the reflux temperature.

5 In another example, sulphur atoms in the compounds, for example when present in a group L¹ or L², may be oxidised to the corresponding sulfoxide or sulphone using an oxidising agent such as a peroxyacid, e.g. 3-chloroperoxybenzoic acid, in an inert solvent such as a halogenated hydrocarbon, e.g. dichloromethane, at around ambient temperature.

In a further example, *N*-oxides of compounds of formula (1) may in general be
10 prepared for example by oxidation of the corresponding nitrogen base as described above in relation to the preparation of intermediates of formula (2).

Salts of compounds of formula (1) may be prepared by reaction of compounds of formula (1) with an appropriate base in a suitable solvent or mixture of solvents, e.g. an organic solvent such as an ether, e.g. diethyl ether, or an alcohol, e.g. ethanol, using
15 conventional procedures.

Where it is desired to obtain a particular enantiomer of a compound of formula (1) this may be produced from a corresponding mixture of enantiomers using any suitable conventional procedure for resolving enantiomers.

Thus, for example, diastereomeric derivatives, e.g. salts, may be produced by
20 reaction of a mixture of enantiomers of formula (1), e.g. a racemate, and an appropriate chiral compound, e.g. a chiral base. The diastereomers may then be separated by any convenient means, for example by crystallisation, and the desired enantiomer recovered, e.g. by treatment with an acid in the instance where the diastereomer is a salt.

In another resolution process a racemate of formula (1) may be separated using
25 chiral High Performance Liquid Chromatography. Alternatively, if desired, a particular enantiomer may be obtained by using an appropriate chiral intermediate in one of the processes described above. Alternatively, a particular enantiomer may be obtained by performing an enantiomer-specific enzymatic biotransformation, e.g. an ester hydrolysis using an esterase, and then purifying only the enantiomerically pure hydrolysed acid from
30 the unreacted ester antipode.

Chromatography, recrystallisation and other conventional separation procedures may also be used with intermediates or final products where it is desired to obtain a particular geometric isomer of the invention.

The following Examples illustrate the invention. All temperatures are in °C. The following abbreviations are used:

EtOAc - ethyl acetate;	MeOH - methanol;
5 DCM - dichloromethane;	EtOH - ethanol; Pyr - pyridine;
DMSO - dimethylsulphoxide;	Me - methyl;
THF - tetrahydrofuran;	h - hour;
MCPBA - 3-chloroperoxybenzoic acid;	
r.t. - room temperature;	DMF - <i>N,N</i> -dimethylformamide;
10 AcOH - acetic acid;	TFAA - trifluoroacetic anhydride;
UHP - urea-hydrogen peroxide complex;	
TLC - thin-layer chromatography;	CDI - 1,1'-carbonyldiimidazole;
EDC - 1-[3-(dimethylamino)propyl]-3-ethylcarbodiimide hydrochloride;	
HOBT - 1-hydroxybenzotriazole hydrate.	

15

All NMRs were obtained either at 300 MHz or 400 MHz.

Compounds were named with the aid of either Beilstein Autonom, supplied by MDL Information Systems GmbH, Theodor-Heuss-Allee 108, D-60486 Frankfurt, Germany, or ACD Labs Name (v. 5.0 or v. 6.0), supplied by Advanced Chemical Development, Toronto, Canada.

LCMS retention times (RT) quoted were generated on a Hewlett Packard 1100 LC/MS using the following method unless otherwise stated: Phenomenex Luna 3 μ C₁₈(2) 50 x 4.6 mm column; mobile phase A = 0.1% formic acid in water; mobile phase B = 0.1% formic acid in MeCN; flow rate of 0.9 mlmin⁻¹; column temperature 40°C.

25

Gradient:

Time (min)	%B
Initial	5
2.0	95
3.0	95
5.0	5
5.5	end

LCMS Method B: Phenomenex Luna 3 μ C₁₈(2) 50 x 4.6 mm column; mobile phase A = 0.1% formic acid in water; mobile phase B = 0.1% formic acid in MeCN; flow rate of 0.9 mlmin⁻¹; column temperature 40°C.

5

Gradient:

Time (min)	%B
Initial	5
2.0	95
4.0	95
5.0	5
5.5	end

LCMS Method C: Varian Polaris 5 μ C₁₈ 50 x 4.6 mm column; mobile phase A = 0.1% formic acid in water; mobile phase B = 0.1% formic acid in MeCN; flow rate of 0.9 mlmin⁻¹; column temperature 40°C.

Gradient:

Time (min)	%B
Initial	5
1.0	95
5.0	95
5.5	end

15

INTERMEDIATE 1

1-Benzenesulfonyl-1*H*-pyrrolo[3,2-*b*]pyridine

Sodium hydride (60% in mineral oil, 203 mg, 5.08 mmol) was washed twice with hexane, and the residue suspended in DMSO (5 ml). 1*H*-Pyrrolo[3,2-*b*]pyridine (500 mg, 4.24 mmol) was added and the reaction stirred for 15 minutes. Benzenesulphonyl chloride (749 mg, 4.42 mmol) was added rapidly and the reaction stirred for 0.5 h. After

quenching with water the product was extracted into EtOAc, dried (Na_2SO_4) and concentrated *in vacuo*. Chromatography (EtOAc, silica) yielded the *title compound* as a white crystalline solid (903 mg, 83%). δ_{H} (d_6 -DMSO) 8.48 (1H, dd, J 0.8, 3.8 Hz), 8.31 (1H, m), 8.15 (1H, d, J 3.8 Hz), 8.04 (2H, m), 7.72 (1H, m), 7.61 (2H, m), 7.36 (1H, dd, J 4.7, 8.4 Hz), 6.98 (1H, dd, J 0.8, 3.8 Hz). LCMS (ES $^+$) RT 3.056 minutes, 259 ($\text{M}+\text{H}$) $^+$.

INTERMEDIATE 2

1-Benzenesulfonyl-1*H*-pyrrolo[3,2-*b*]pyridine 4-oxide

10 Intermediate 1 (903 mg, 3.5 mmol) was dissolved in EtOAc (6 ml) and treated with MCPBA (903 mg, 5.25 mmol). After stirring for 24 h the reaction mixture was washed with saturated sodium hydrogencarbonate solution, dried (MgSO_4) and concentrated *in vacuo*. Chromatography (EtOAc, silica) yielded the *title compound* as a white foam (860 mg, 90%). δ_{H} (d_6 -DMSO) 8.24 (1H, d, J 6.6 Hz), 8.15 (3H, m), 7.9 (1H, d, J 8.3 Hz), 7.76 (1H, m), 7.68 (2H, m), 7.35 (1H, dd, J 6.3, 8.5 Hz), 7.1 (1H, dd, J 0.8, 3.8 Hz). LCMS (ES $^+$) RT 2.639 minutes, 275 ($\text{M}+\text{H}$) $^+$.

INTERMEDIATE 3

20 1-Benzenesulfonyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 2 (0.86 g, 3.1 mmol) was dissolved in DMF (8 ml) and treated with TFAA (6.6 g, 31.4 mmol) at r.t. and stirred for 4 days. The reaction mixture was concentrated *in vacuo*, redissolved in EtOH, concentrated again, then partitioned between water and toluene. The organic layer was dried (MgSO_4) and concentrated to yield the 25 *title compound* as a beige solid (657 mg, 77%). δ_{H} (d_6 -DMSO) 8.20 (1H, d, J 8.6 Hz), 8.11 (2H, m), 7.94 (1H, d, J 3.7 Hz), 7.88 (1H, d, J 8.4 Hz), 7.78 (2H, m), 6.58 (1H, d, J 3.6 Hz), 6.45 (1H, d, J 9.6 Hz). LCMS (ES $^+$) RT 2.775 minutes, 275 ($\text{M}+\text{H}$) $^+$.

INTERMEDIATE 4

30

1-Benzenesulfonyl-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 3 (300 mg, 1.09 mmol) was combined with phenylboronic acid (343 mg, 2.19 mmol), copper(II) acetate (437 mg, 2.19 mmol), and pyridine (319 mg, 0.41 ml,

3.28 mmol) in DCM. The reaction was stirred at r.t. for 24 h then diluted with further DCM, washed with water, and the organic layer dried and concentrated *in vacuo*. Column chromatography (EtOAc, silica) yielded the *title compound* (150 mg, 39%). δ_H (d_6 -DMSO) 7.95 (1H, d, J 9.8 Hz), 7.90 (2H, d, J 8.5 Hz), 7.60 (2H, m), 7.53 (2H, m), 5 7.35 (3H, m), 7.16 (2H, d, J 1.7, 8.3 Hz), 6.23 (1H, d, J 9.8 Hz), 5.70 (1H, dd, J 0.6, 3.7 Hz). LCMS (ES⁺) RT 3.312 minutes, 351 (M+H)⁺.

INTERMEDIATE 5

10 4-Phenyl-1,4-dihydro-5H-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 4 (1.5 g, 4.3 mmol) in MeOH (10 ml) was treated with 2M sodium hydroxide (5 ml) and stirred at r.t. for 2 h. The organic phase was removed under reduced pressure, the residue neutralised with 2M hydrochloric acid then extracted into DCM, dried ($MgSO_4$) and concentrated *in vacuo* to yield the *title compound* (765 mg, 85%). δ_H ($CDCl_3$) 9.30 (1H, br s), 7.44 (3H, m), 7.36 (3H, m), 6.84 (1H, t, J 3.0 Hz), 6.37 (1H, d, J 9.4 Hz), 5.59 (1H, t, J 2.3 Hz).

INTERMEDIATE 6

20 1-Benzenesulfonyl-4-(4-methoxyphenyl)-1,4-dihydro-5H-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 3 (300 mg, 1.09 mmol) and 4-methoxyphenylboronic acid (333 mg, 2.19 mmol) by the method of Intermediate 4 to give the *title compound* (110 mg, 27%). δ_H (d_6 -DMSO) 8.12 (1H, d, J 9.8 Hz), 8.06 (2H, m), 7.75 (4H, m), 7.25 (2H, dd, J 2.2, 9.0 Hz), 7.05 (2H, dd, J 2.2, 9.0 Hz), 6.40 (1H, d, J 9.8 Hz), 5.91 (1H, dd, J 0.5, 3.6 Hz), 3.81 (3H, s). LCMS (ES⁺) RT 3.309 minutes, 381 (M+H)⁺.

INTERMEDIATE 7

4-(4-Methoxyphenyl)-1,4-dihydro-5H-pyrrolo[3,2-*b*]pyridin-5-one

30 From Intermediate 6 (25 mg, 0.066 mmol) by the method of Intermediate 5 to give the *title compound* (16 mg, 100%). δ_H ($CDCl_3$) 9.04 (1H, br s), 7.46 (1H, dd, J 0.5, 9.3 Hz), 7.25 (2H, m), 6.96 (2H, m), 6.89 (1H, apparent t, J 2.9 Hz), 6.38 (1H, d, J 9.3 Hz), 5.63 (1H, apparent t, J 2.3 Hz), 3.78 (3H, s). HPLC RT 2.608 minutes.

INTERMEDIATE 8(Phenyl)(pyrrolo[3,2-*b*]pyridin-1-yl)methanone

5 By the method of Intermediate 1 using benzoyl chloride (596 mg, 4.42 mmol) in place of benzenesulphonyl chloride. Yellow crystalline solid (429 mg, 44%). δ_H (d_6 -DMSO) 8.60 (1H, m), 8.50 (1H, m), 7.79 (2H, m), 7.74 (2H, m), 7.63 (2H, m), 7.39 (1H, dd, J 4.7, 8.3 Hz), 6.86 (1H, dd, J 0.6, 3.9 Hz). LCMS (ES⁺) RT 2.622 minutes, 223 (M+H)⁺.

10

INTERMEDIATE 9(4-Oxypyrrrolo[3,2-*b*]pyridin-1-yl)(phenyl)methanone

From Intermediate 8 (429 mg, 1.93 mmol) by the method of Intermediate 2.

15 White foam (403 mg, 88%). δ_H (d_6 -DMSO) 8.28 (1H, d, J 7.7 Hz), 8.10 (1H, d, J 8.8 Hz), 7.81 (2H, d, J 7.7 Hz), 7.74 (1H, m), 7.67 (3H, m), 7.38 (1H, dd, J 6.4, 8.5 Hz), 6.98 (1H, m). LCMS (ES⁺) RT 2.612 minutes, 239 (M+H)⁺.

INTERMEDIATE 10

20

1-Benzoyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 9 (400 mg, 1.68 mmol) was dissolved in DMF (4 mL) and treated dropwise with TFAA (3.5 g, 16.8 mmol) then stirred at r.t. for 4 days. The reaction mixture was concentrated *in vacuo*, redissolved in ethanol, concentrated again, and finally 25 azeotroped with heptane to remove the remaining TFAA. The residue was partitioned between toluene and water, and the organic phase separated, dried (Na₂SO₄) and concentrated *in vacuo* to give the *title compound* as a white crystalline solid (218 mg, 55%). δ_H (d_6 -DMSO) 8.18 (1H, d, J 9.6 Hz), 7.78-7.64 (3H, m), 7.64-7.51 (2H, m), 7.31 (1H, d, J 3.5 Hz), 6.30 (1H, m), 6.25 (1H, d, J 9.7 Hz). HPLC RT 2.717 minutes.

30

INTERMEDIATE 114-Phenyl-1-(phenylsulfonyl)-2-(pyrrolidin-1-ylsulfonyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

5 A solution of Intermediate 4 (400 mg, 1.14 mmol) in THF (6 ml) was cooled to -78°C under nitrogen and treated dropwise with *n*-BuLi (1.6 M in hexanes, 0.86 ml, 1.37 mmol) and stirred for 30 minutes. Sulphur dioxide was blown through the reaction mixture at -78°C for 5 minutes and the reaction mixture was then allowed to warm to r.t. The solvents were removed *in vacuo* and the residue suspended in DCM (10ml). *N*-Chlorosuccinimide (203 mg, 1.14 mmol) was added and the reaction stirred for 30 minutes, then treated with pyrrolidine (81 mg, 0.095 ml, 1.14 mmol). After stirring for a further 30 minutes the reaction was partitioned between water and DCM, and the organic phase was dried (Na_2SO_4), concentrated and purified by chromatography (EtOAc, silica) to give the *title compound* (210 mg, 38%). LCMS Method B (ES^+) RT 2.410 minutes, 484 ($\text{M}+\text{H}^+$).

10 15

INTERMEDIATE 124-Phenyl-2-(pyrrolidin-1-ylsulfonyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 11 (200 mg, 0.41 mmol) was suspended in 2M sodium hydroxide (10 ml) and treated with sufficient MeOH to solubilise. The reaction was stirred at r.t. for 3 days. The MeOH was removed *in vacuo* and the residual aqueous solution washed with DCM. After acidification with conc. HCl the product was extracted into DCM, and the organic phase dried (Na_2SO_4) and concentrated *in vacuo*. Purification of the residue by chromatography (EtOAc, silica) gave the *title compound* (140 mg, 100%). δ_{H} (d_6 -DMSO)

25 7.76 (1H, d, *J* 9.6 Hz), 7.67-7.61 (2H, m), 7.58-7.55 (1H, m), 7.48-7.45 (2H, m), 6.51 (1H, d, *J* 9.6 Hz), 5.97 (1H, s), 3.22-3.17 (4H, m), 1.74-1.69 (4H, m). LCMS (ES^+) RT 2.801 minutes, 344 ($\text{M}+\text{H}^+$).

INTERMEDIATE 13

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Ethyl (2*Z*)-2-hydroxy-3-(3-nitropyridin-2-yl)acrylate

Sodium metal (400 mg, 17.4 mmol) was added portionwise to EtOH (50 ml), followed by diethyl oxalate (2.4 ml, 17.4 mmol), and stirred at r.t. for 5 min. 2-Methyl-3-

nitropyridine (2.00 g, 14.6 mmol) was added, giving a deep purple solution that changed over time to red-orange. After stirring for 3 days at r.t. a red precipitate had formed which was collected by filtration and washed with ether (4 x 20 ml). The solid residue was suspended in water and the suspension acidified to pH 4 with AcOH. The resulting
5 orange precipitate was collected by filtration, washed with ethanol (10 ml) and dried *in vacuo* to give the *title compound* (1.17 g, 34%). δ_H ($CDCl_3$) 8.77-8.75 (1H, m), 8.54-8.50 (1H, m), 7.47-7.43 (2H, m), 4.48 (2H, q, J 7.1 Hz), 1.49 (3H, t, J 7.1 Hz). LCMS (ES⁺) RT 3.32 minutes, 239 ($M+H^+$).

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INTERMEDIATE 14Ethyl 1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

A mixture of Intermediate 13 (6.30 g, 26.5 mmol) and iron powder (325 mesh, 13.3 g, 238.2 mmol) in EtOH (100 ml) and AcOH (100 ml) was heated at reflux for 2 h.
15 Volatiles were removed *in vacuo* and toluene used to azeotrope excess AcOH (3 x 100 ml). The brown residue was suspended in EtOAc and passed through a plug of silica gel, eluting the desired product with EtOAc. Concentration of the eluent *in vacuo* gave the *title compound* as an off-white solid (4.32 g, 86%). δ_H (d_3 -MeOD) 8.33-8.30 (1H, m), 7.84-7.80 (1H, m), 7.22 (1H, dd, J 4.6, 8.4 Hz), 7.14 (1H, d, J 0.9 Hz), 4.33 (2H, q, J 7.1 Hz), 1.33 (3H, t, J 7.1 Hz). LCMS Method C (ES⁺) RT 1.22 minutes, 191 ($M+H^+$).
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INTERMEDIATE 15Ethyl 1-benzyl-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

25 A solution of Intermediate 14 (200 mg, 1.05 mmol) in DMF (10 ml) was treated with sodium hydride (60% in mineral oil, 42 mg, 1.05 mmol) at r.t. until effervescence ceased. After cooling to 0°C, benzyl bromide (0.13 ml, 1.05 mmol) was added and the reaction stirred for 10 minutes. The reaction mixture was partitioned between water and EtOAc (20 ml each) and extracted with EtOAc (20 ml). The combined organics were
30 washed with brine (20 ml), dried ($MgSO_4$), filtered and concentrated *in vacuo*. Purification by chromatography (5%-20% MeOH in DCM, silica) gave the *title compound* as a yellow solid (143 mg, 48%). δ_H ($CDCl_3$) 8.38-8.36 (1H, m), 7.50-7.46

(1H, m), 7.36 (1H, s), 7.20-7.01 (4H, m), 6.88-6.85 (2H, m), 5.69 (2H, s), 4.20 (2H, q, *J* 7.1 Hz), 1.21 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.16 minutes, 281 (M+H⁺).

INTERMEDIATE 16

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Ethyl 1-benzyl-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate 4-oxide

A solution of Intermediate 15 (140 mg, 0.5 mmol) in THF (1 ml) at 0°C was treated with UHP (99 mg, 1.05 mmol). Slow addition of TFAA (0.15 ml, 1.05 mmol) was followed by stirring at 0°C for 30 min. The reaction mixture was treated with dilute aqueous sodium thiosulfate solution (5 ml) for 30 min then acidified with dilute HCl. Extraction with DCM (2 x 5 ml), drying (MgSO₄), filtration and concentration *in vacuo*, then purification by chromatography (50% EtOAc in DCM-15% MeOH in EtOAc, silica) gave the *title compound* as a yellow solid (120 mg, 81%). δ_H (CDCl₃) 8.14 (1H, d, *J* 8.5 Hz), 7.68 (1H, s), 7.15-7.12 (4H, m), 7.08-6.95 (3H, m), 5.79 (2H, s), 4.29 (2H, q, *J* 7.1 Hz), 1.30 (3H, t, *J* 7.1 Hz). LCMS Method C (ES⁺) RT 2.33 minutes, 297 (M+H⁺).

INTERMEDIATE 17

20 A solution of Intermediate 16 (120 mg, 0.40 mmol) in DMF (1 ml) was treated with TFAA (0.57 ml, 4.05 mmol) and stirred at r.t. for 2 h. The reaction mixture was partitioned between EtOAc and NaHCO₃ (20 ml each) and the organics washed with water (5 ml), then brine (5 ml), dried (MgSO₄), filtered and concentrated *in vacuo* to give the *title compound*. δ_H (CDCl₃) 7.47 (1H, d, *J* 9.6 Hz), 7.16-7.25 (3H, m), 6.97-6.95 (3H, m), 6.46 (1H, d, *J* 9.6 Hz), 5.71 (2H, s), 4.22 (2H, q, *J* 7.1 Hz), 1.26 (3H, t, *J* 7.1 Hz). LCMS Method C (ES⁺) RT 2.25 minutes, 297 (M+H⁺).

INTERMEDIATE 18

30 Ethyl 1-(3-chloro-4-fluorobenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 14 (1.00 g, 5.26 mmol) and 3-chloro-4-fluorobenzyl bromide (1.30 g, 5.80 mmol) by the method of Intermediate 15. Purification by chromatography (20%-50% EtOAc in hexane, silica) gave the *title compound* (1.09 g, 62%). δ_H (CDCl₃)

8.64-8.63 (1H, m), 7.67 (1H, d, *J* 9.6 Hz), 7.57 (1H, s), 7.29-7.25 (1H, m), 7.16-7.04 (2H, m), 6.95-6.90 (1H, m), 5.81 (2H, s), 4.42 (2H, q, *J* 7.1 Hz), 1.43 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.36 minutes, 333 [³⁵Cl]:335 [³⁷Cl] 3:1 (M+H⁺).

5

INTERMEDIATE 19Ethyl 1-(3-chloro-4-fluorobenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate 4-oxide

From Intermediate 18 (1.09 g, 3.28 mmol), with UHP (650 mg, 6.89 mmol) and TFAA (0.97 ml, 6.89 mmol) by the method of Intermediate 16. Purification by chromatography (5%-15% MeOH in EtOAc, silica) gave the *title compound* (920 mg, 81%). δ_H (CDCl₃) 8.14-8.11 (1H, m), 7.68 (1H, s), 7.18-7.14 (1H, m), 7.04-6.90 (3H, m), 6.81-6.76 (1H, m), 5.65 (2H, s), 4.24 (2H, q, *J* 7.1 Hz), 1.26 (3H, t, *J* 7.1 Hz). LCMS Method C (ES⁺) RT 2.39 minutes, 349 [³⁵Cl]:351 [³⁷Cl] 3:1 (M+H⁺).

15

INTERMEDIATE 20Ethyl 1-(3-chloro-4-fluorobenzyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 19 (910 mg, 2.61 mmol) and TFAA (3.70 ml, 26.1 mmol) by the method of Intermediate 17. *Title compound* required no further purification (886 mg, 97%). δ_H (CDCl₃) 7.84 (1H, d, *J* 9.6 Hz), 7.11-6.68 (3H, m), 6.82 (1H, s), 6.42 (1H, d, *J* 9.6 Hz), 5.69 (2H, s), 4.23 (2H, q, *J* 7.1 Hz), 1.23 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.22 minutes, 349 [³⁵Cl]:351 [³⁷Cl] 3:1 (M+H⁺).

25

INTERMEDIATE 21Ethyl 1-(3-methylbenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 14 (1.00 g, 5.26 mmol) and 3-methylbenzyl bromide (0.78 ml, 5.80 mmol) according to the procedure for Intermediate 15. Purification by chromatography (50% EtOAc in hexane, silica) gave the *title compound* (520 mg, 34%). δ_H (CDCl₃) 8.38-8.36 (1H, m), 7.45 (1H, d, *J* 9.6 Hz), 7.32 (1H, s), 7.03-6.82 (3H, m), 6.72-6.60 (2H, m), 5.63 (2H, s), 4.18 (2H, q, *J* 7.1 Hz), 2.07 (3H, s), 1.19 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.16 minutes, 294 (M+H⁺).

INTERMEDIATE 22Ethyl 1-(3-methylbenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate 4-oxide

5 From Intermediate 21 (520 mg, 1.77 mmol), using UHP (350 mg, 3.71 mmol) and TFAA (0.53 ml, 3.71 mmol) according to the procedure for Intermediate 16. Purification by chromatography (2%-15% MeOH in EtOAc, silica) gave the *title compound* (455 mg, 83%). δ_H ($CDCl_3$) 8.18-8.14 (1H, m), 7.73 (1H, s), 7.28-7.21 (1H, m), 7.14-6.98 (3H, m), 6.76-6.73 (2H, m), 5.75 (2H, s), 4.29 (2H, q, J 7.1 Hz), 2.20 (3H, s), 1.31 (3H, t, J 7.1 Hz).

10 LCMS (ES⁺) RT 2.36 minutes, 311 (M+H⁺).

INTERMEDIATE 23Ethyl 1-(3-methylbenzyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

15 From Intermediate 22 (450 mg, 1.45 mmol) using TFAA (2.05 ml, 14.5 mmol) according to the procedure for Intermediate 17. *Title compound* required no further purification (354 mg, 79%). δ_H ($CDCl_3$) 7.80 (1H, d, J 9.6 Hz), 7.12-7.09 (1H, m), 7.03-7.00 (1H, m), 6.84 (2H, s), 6.78-6.76 (1H, m), 6.41 (1H, d, J 9.6 Hz), 5.76 (2H, s), 4.29 (2H, q, J 7.1 Hz), 2.23 (3H, s), 1.29 (3H, t, J 7.1 Hz). LCMS (ES⁺) RT 3.14 minutes, 311 (M+H⁺).

INTERMEDIATE 24Ethyl 1-(3-chlorobenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

25 From Intermediate 14 (1.00 g, 5.26 mmol), using 3-chlorobenzyl bromide (0.76 ml, 5.80 mmol) according to the procedure for Intermediate 15. Purification by chromatography (30% EtOAc in hexane, silica) gave the *title compound* (833 mg, 50%). δ_H ($CDCl_3$) 8.50-8.47 (1H, m), 7.65 (1H, m), 7.45 (1H, s), 7.14-7.06 (3H, m), 6.91-6.90 (1H, m), 6.78-6.75 (1H, m), 5.69 (2H, s), 4.24 (2H, q, J 7.1 Hz), 1.26 (3H, t, J 7.1 Hz).

30 LCMS (ES⁺) RT 3.29 minutes, 315 [^{35}Cl]:317 [^{37}Cl] 3:1(M+H⁺).

INTERMEDIATE 25Ethyl 1-(3-chlorobenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate 4-oxide

From Intermediate 24 (830 mg, 2.64 mmol), using UHP (522 mg, 5.55 mol) and TFAA (0.79 ml, 5.55 mmol) according to the procedure for Intermediate 16. Purification by chromatography (2%-10% MeOH in EtOAc, silica) gave the *title compound* (309 mg, 35%). δ_H ($CDCl_3$) 8.17-8.14 (1H, m), 7.69 (1H, s), 7.21-7.07 (4H, m), 6.96-6.82 (2H, m), 5.75 (2H, s), 4.29 (2H, q, J 7.1 Hz), 1.31 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 2.38 minutes, 331 [^{35}Cl]:333 [^{37}Cl] 3:1 (M+H $^+$).

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INTERMEDIATE 26Ethyl 1-(3-chlorobenzyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 25 (300 mg, 0.91 mmol) using TFAA (1.30 ml, 9.10 mmol) according to the procedure for Intermediate 17. The *title compound* required no further purification (372 mg, 100%). δ_H ($CDCl_3$) 7.25-7.15 (2H, m), 7.16 (1H, d, J 9.6 Hz), 6.96 (1H, s), 6.86-6.80 (2H, m), 6.46 (1H, d, J 9.6 Hz), 5.67 (2H, s), 4.24 (2H, q, J 7.1 Hz), 1.28 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 3.19 minutes, 331 [^{35}Cl]:333 [^{37}Cl] 3:1 (M+H $^+$).

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INTERMEDIATE 271-(3-Chloro-4-fluorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

Example 24 (104 mg, 0.24 mmol) was treated with 1M sodium hydroxide (0.24 ml, 0.24 mmol) in EtOH (0.5 ml) under reflux for 4 hours. EtOH was removed *in vacuo* and the residue treated with 2M HCl, producing a white precipitate. This was collected by filtration, washed with water (1 ml) and Et₂O (1 ml), and dried *in vacuo* to give the *title compound* (80 mg, 82%) which required no further purification. δ_H (d_6 -DMSO) 8.09 (1H, d, J 9.7 Hz), 7.64-7.42 (7H, m), 7.17-7.12 (1H, m), 6.55 (1H, d, J 9.7 Hz), 6.21 (1H, s), 5.88 (2H, s). LCMS (ES^+) RT 3.11 minutes, 397 [^{35}Cl]:399 [^{37}Cl] 3:1 (M+H $^+$).

INTERMEDIATE 281-(3-Methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

5 Example 25 (261 mg, 0.67 mmol) was treated with lithium hydroxide (85 mg, 2.02 mmol) in EtOH and water (2 ml each) at r.t. overnight, followed by reflux for 4 hours. EtOH was removed *in vacuo* and the residue treated with 2M HCl, producing a white precipitate. This was collected by filtration, washed with water (2 ml) and Et₂O (4 ml), and dried *in vacuo* to give the *title compound* (228 mg, 94%) which required no further purification. δ_H (d₆-DMSO) 7.94 (1H, d, *J* 9.7 Hz), 7.60-7.50 (3H, m), 7.47-7.39 (2H, m), 7.19 (1H, t, *J* 7.5 Hz), 7.06 (1H, d, *J* 8.7 Hz), 7.00 (1H, s), 6.86 (1H, d, *J* 7.3 Hz), 6.42 (1H, d, *J* 9.7 Hz), 6.09 (1H, s), 5.81 (2H, s), 2.26 (3H, s). LCMS (ES⁺) RT 3.01 minutes, 359 (M+H⁺).

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INTERMEDIATE 291-(3-Chlorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

Example 26 (310 mg, 0.76 mmol) was treated with lithium hydroxide (96 mg, 2.29 mmol) in EtOH and water (2 ml each) at r.t. overnight. EtOH was removed *in vacuo* and the residue treated with 2M HCl, producing a white precipitate. This was collected by filtration, washed with water (2 ml) and Et₂O (4 ml), and dried *in vacuo* to give the *title compound* (258 mg, 89%) which required no further purification. δ_H (d₆-DMSO) 8.22 (1H, d, *J* 9.7 Hz), 7.83-7.54 (7H, m), 7.46 (1H, s), 7.26-7.24 (1H, m), 6.68 (1H, d, *J* 9.7 Hz), 6.35 (1H, s), 6.06 (2H, s). LCMS (ES⁺) RT 3.08 minutes, 379 [³⁵Cl]:381 [³⁷Cl] 3:1 (M+H⁺).

INTERMEDIATE 30

30 1-(3-Chlorobenzyl)-4-(1*H*-indol-5-yl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

1M sodium hydroxide solution (0.75 ml, 0.75 mmol) was added to a suspension of Example 39 (243 mg, 0.55 mmol) in EtOH and water (2 ml each) and heated at reflux

overnight. The reaction mixture was concentrated *in vacuo* giving the *title compound* (245 mg, 100%), which was used without further purification. δ_H (d_6 -DMSO) 7.54 (1H, d, J 9.5 Hz), 7.46-7.37 (3H, m), 7.28-7.12 (4H, m), 6.90 (1H, dd, J 8.5, 2.0 Hz), 6.42 (1H, d, J 3.0 Hz), 6.05 (1H, d, J 9.5 Hz), 5.96 (1H, s), 5.67 (2H, s). LCMS (ES $^+$) RT 3.02
5 minutes, 418 [^{35}Cl]:420 [^{37}Cl] 3:1 (M+H $^+$).

INTERMEDIATE 31

Ethyl 1-(4-fluoro-3-methylbenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

10 From Intermediate 14 (1.0 g, 5.26 mmol) using 4-fluoro-3-methylbenzyl bromide (1.18 g, 5.80 mmol) according to the procedure for Intermediate 15. Purification by chromatography (5%-10% EtOAc in DCM, silica) gave the *title compound* (670 mg, 40%). δ_H (CDCl $_3$) 8.61 (1H, d, J 4.4 Hz), 7.68 (1H, d, J 8.5 Hz), 7.55 (1H, s), 7.24 (1H, dd, J 4.4, 8.5 Hz), 6.96-6.81 (3H, m), 5.79 (2H, s), 4.39 (2H, q, J 7.1 Hz), 2.20 (3H, d, J 1.6 Hz), 1.41 (3H, t, J 7.1 Hz). LCMS (ES $^+$) RT 3.20 minutes, 313 (M+H $^+$).
15

INTERMEDIATE 32

Ethyl 1-(4-fluoro-3-methylbenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate 4-oxide

20 From Intermediate 31 (630 mg, 2.02 mmol) according to the procedure for Intermediate 16. Purification by chromatography (10% MeOH in EtOAc, silica) gave the *title compound* (680 mg, 100%). δ_H (CDCl $_3$) 8.23 (1H, d, J 6.0 Hz), 7.45 (1H, s), 7.32 (1H, d, J 8.5 Hz), 7.17 (1H, dd, J 6.0, 8.5 Hz), 6.97-6.84 (3H, m), 5.81 (2H, s), 4.40 (2H, q, J 7.1 Hz), 2.23 (3H, d, J 1.6 Hz), 1.41 (3H, t, J 7.1 Hz). LCMS (ES $^+$) RT 3.13 minutes,
25 329 (M+H $^+$).

INTERMEDIATE 33

Ethyl 1-(4-fluoro-3-methylbenzyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-

30 carboxylate

From Intermediate 32 (680 mg, 2.07 mmol), according to the procedure for Intermediate 17. The *title compound* required no further purification (680 mg, 100%). δ_H (CDCl $_3$) 8.16 (1H, d, J 9.7 Hz), 7.26-7.18 (2H, m), 7.04-6.99 (1H, m), 6.83 (1H, s), 6.47

(1H, d, *J* 9.7 Hz), 5.87 (2H, s), 4.42 (2H, q, *J* 7.1 Hz), 2.33 (3H, s), 1.43 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.29 minutes, 329 (M+H⁺).

INTERMEDIATE 34

5

1-(Phenylsulfonyl)-4-(3-thienyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 3 (5.0 g, 18.2 mmol) using 3-thiopheneboronic acid (5 g, 36.5 mmol) according to the procedure for Example 23. Purification by chromatography (5%-20% EtOAc in DCM, silica) gave the *title compound* (1.23 g, 19%). δ_H (d_3 -MeOD) 8.28 (1H, d, *J* 9.7 Hz), 8.00 (2H, d, *J* 8.0 Hz), 7.74-7.68 (2H, m), 7.62-7.55 (4H, m), 7.11-7.08 (1H, m), 6.51 (1H, d, *J* 9.7 Hz), 6.09 (1H, d, *J* 3.6 Hz). LCMS (ES⁺) RT 3.22 minutes, 357 (M+H⁺).

INTERMEDIATE 35

15

4-(3-Thienyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 34 (1.20 g, 3.37 mmol), according to the method of Intermediate 5. The *title compound* required no further purification (680 mg, 94%). δ_H (d_3 -MeOD) 8.02 (1H, d, *J* 9.3 Hz), 7.87-7.82 (2H, m), 7.44-7.40 (2H, m), 6.61 (1H, d, *J* 9.3 Hz), 6.07-6.06 (1H, m). LCMS (ES⁺) RT 2.44 minutes, 217 (M+H⁺).

INTERMEDIATE 36

Ethyl 1-(2-cyanobenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

25 From Intermediate 14 (1.0 g, 5.26 mmol) using 2-cyanobenzyl bromide (1.14 g, 5.80 mmol) according to the procedure for Intermediate 15. Purification by chromatography (5%-10% EtOAc in DCM, silica) gave the *title compound* (600 mg, 37%). δ_H ($CDCl_3$) 8.79 (1H, br s), 7.90-7.87 (1H, m), 7.79-7.76 (2H, m), 7.58-7.39 (3H, m), 6.73 (1H, d, *J* 7.6 Hz), 6.23 (2H, s), 4.53 (2H, q, *J* 7.1 Hz), 1.54 (3H, t, *J* 7.1 Hz).
30 LCMS (ES⁺) RT 2.92 minutes, 306 (M+H⁺).

INTERMEDIATE 37Ethyl 1-(2-cyanobenzyl)-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate 4-oxide

From Intermediate 36 (600 mg, 1.97 mmol) according to the method of

5 Intermediate 16. Purification by chromatography (5%-10% MeOH in EtOAc, silica) gave the *title compound* (600 mg, 100%). δ_H ($CDCl_3$) 8.02 (1H, d, J 6.0 Hz), 7.62 (1H, s), 7.54 (1H, d, J 7.5 Hz), 7.28-7.16 (2H, m), 7.04-6.94 (2H, m), 6.42 (1H, d, J 7.5 Hz), 5.88 (2H, s), 4.16 (2H, q, J 7.1 Hz), 1.17 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 2.81 minutes, 322 ($M+H^+$).

10

INTERMEDIATE 38Ethyl 1-(2-cyanobenzyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 37 (680 mg, 2.07 mmol) according to the procedure for

15 Intermediate 17. The *title compound* required no further purification (680 mg, 100%). δ_H ($CDCl_3$) 7.74-7.81 (2H, m), 7.41 (1H, dt, J 7.7, 1.3 Hz), 7.27-7.32 (1H, m), 6.58 (1H, s), 6.27 (1H, d, J 7.7 Hz), 6.18 (1H, d, J 9.7 Hz), 5.81 (2H, s), 4.02 (2H, q, J 7.1 Hz), 1.04 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 2.88 minutes, 322 ($M+H^+$).

20

INTERMEDIATE 39Ethyl 1-(2-cyanobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 38 (630 mg, 1.96 mmol), using phenylboronic acid (488 mg,

25 4.00 mmol), according to the procedure for Example 23. Purification by chromatography (10%-50% EtOAc in DCM, silica) gave the *title compound* (370 mg, 47%). δ_H (d_3 -MeOD) 7.88 (1H, d, J 9.7 Hz), 7.75 (1H, dd, J 1.3, 7.6 Hz), 7.62-7.46 (4H, m), 7.40-7.36 (3H, m), 6.61 (1H, d, J 7.8 Hz), 6.55 (1H, d, J 9.7 Hz), 6.31 (1H, s), 6.01 (2H, s), 4.14 (2H, q, J 7.1 Hz), 1.14 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 3.41 minutes, 398 ($M+H^+$).

30

INTERMEDIATE 40

1-(2-Cyanobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

5 From Intermediate 39 (337 mg, 0.85 mmol) according to the procedure of Intermediate 30. δ_H (d_6 -DMSO) 7.84 (1H, dd, J 1.1, 7.7 Hz), 7.65 (1H, d, J 9.5 Hz), 7.60-7.36 (7H, m), 6.47 (1H, d, J 7.7 Hz), 6.26 (2H, s), 6.14 (1H, d, J 9.5 Hz), 5.84 (1H, s). LCMS (ES $^+$) RT 2.80 minutes, 370 (M+H $^+$).

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INTERMEDIATE 41

1-(4-Fluoro-3-methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

From Example 42 (541 mg, 1.34 mmol) according to the procedure of
15 Intermediate 30. The *title compound* required no further purification (360 mg, 100%). δ_H (d_6 -DMSO) 7.76 (1H, d, J 9.5 Hz), 7.65-7.50 (3H, m), 7.42-7.40 (2H, m), 7.28 (1H, d, J 7.7 Hz), 7.17-7.06 (2H, m), 6.19 (1H, d, J 9.5 Hz), 6.03 (2H, s), 5.82 (1H, s), 2.23 (3H, s). LCMS (ES $^+$) RT 3.04 minutes, 377 (M+H $^+$).

20

INTERMEDIATE 42

1-(3-Chlorobenzyl)-4-(4-methylphenyl)-5-oxo-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxylic acid

From Example 44 (300 mg, 0.71 mmol) according to the procedure of
25 Intermediate 30. The *title compound* required no further purification (315 mg, 100%). δ_H (d_6 -DMSO) 7.95 (1H, d, J 9.5 Hz), 7.59-7.40 (8H, m), 6.48 (1H, d, J 9.5 Hz), 6.25 (2H, s), 5.99 (1H, s), 2.62 (3H, s). LCMS (ES $^+$) RT 3.16 minutes, 393 [^{35}Cl]:395 [^{37}Cl] 3:1 (M+H $^+$).

30

INTERMEDIATE 431-Benzyl-1*H*-pyrrolo[3,2-*b*]pyridine

A solution of 1*H*-pyrrolo[3,2-*b*]pyridine (1.30 g, 9.42 mmol) in DMF (15 ml) was
5 treated with sodium hydride (60% in mineral oil, 434 mg, 11.3 mmol) at r.t. until
effervescence ceased. After cooling to 0°C benzyl bromide (1.68 ml, 14.1 mmol) was
added and the reaction warmed at 60°C overnight. The reaction mixture was partitioned
between water and EtOAc (50 ml each) and extracted with EtOAc (50 ml). The
combined organics were washed with brine (20 ml), dried (MgSO_4), filtered and
10 concentrated *in vacuo* to give the *title compound* as a brown oil (1.20 g, 58%). δ_{H}
(CDCl_3) 8.39 (1H, d, J 4.4 Hz), 7.48 (1H, d, J 8.2 Hz), 7.30 (1H, d, J 3.1 Hz), 7.26-7.18
(3H, m), 7.04-6.69 (3H, m), 6.67 (1H, d, J 4.4 Hz), 5.11 (2H, s). LCMS (ES⁺) RT 1.56
min, 209 (M+H⁺).

15

INTERMEDIATE 441-Benzyl-1*H*-pyrrolo[3,2-*b*]pyridine 4-oxide

From Intermediate 43 (1.20 g, 5.50 mmol) according to the procedure for
Intermediate 9. The *title compound* required no further purification (1.23 g, 100%). δ_{H}
20 (CDCl_3) 8.10 (1H, d, J 6.0 Hz), 7.22-7.27 (5H, m), 7.10-7.09 (2H, m), 7.01-6.99 (2H, m),
5.25 (2H, s). LCMS (ES⁺) RT 2.61 min, 225 (M+H⁺).

INTERMEDIATE 4525 1-Benzyl-3-(trifluoroacetyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

A solution of Intermediate 44 (1.23 g, 5.50 mmol) in DMF (20 ml) was treated
with TFAA (7.75 ml, 55.0 mmol) at 0°C. The reaction mixture was stirred at r.t. for 3 h.
The reaction mixture was concentrated *in vacuo*, with use of a hot (approx 60°C) water-
bath. Purification by chromatography (80% EtOAc in hexane to neat EtOAc, silica) gave
30 the *title compound* (560 mg, 45%). δ_{H} (CDCl_3) 7.88 (1H, s), 7.52 (1H, d, J 2.0 Hz), 7.30-
7.25 (3H, m), 7.05-7.03 (2H, m), 6.36 (1H, d, J 9.0 Hz), 5.16 (2H, s). LCMS (ES⁺) RT
3.12 min, 321 (M+H⁺).

EXAMPLE 11-Benzyl-4-phenyl-1,4-dihydro-5H-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 5 (140 mg, 0.67 mmol) was added to sodium hydride (60% suspension in mineral oil, 53 mg, 1.33 mmol) suspended in THF (2 ml). Benzyl chloride (125 mg, 0.73 mmol) was added and the reaction stirred at r.t. for 30 minutes. The reaction mixture was concentrated *in vacuo* and purification by chromatography (gradient elution DCM-EtOAc, silica) gave the *title compound* (168 mg, 84%). δ_H (CDCl₃) 7.56 (2H, m), 7.46 (4H, m), 7.38 (3H, m), 7.17 (2H, m), 6.92 (1H, d, *J* 3.0 Hz), 6.44 (1H, d, *J* 9.5 Hz), 5.68 (1H, d, *J* 2.7 Hz), 5.25 (2H, s). LCMS (ES⁺) RT 3.241 minutes, 301 (M+H)⁺.

EXAMPLES 2 TO 5

15 The compounds of Examples 2-5 were made by a similar procedure to the compound of Example 1, using Intermediate 5 and the appropriate substituted benzyl chloride.

EXAMPLE 2

20

1-(3-Chlorobenzyl)-4-phenyl-1,4-dihydro-5H-pyrrolo[3,2-*b*]pyridin-5-one

δ_H (CDCl₃) 7.49 (2H, m), 7.36 (4H, m), 7.23 (2H, m), 7.06 (1H, d, *J* 0.7 Hz), 6.94 (1H, m), 6.84 (1H, d, *J* 3.0 Hz), 6.36 (1H, d, *J* 9.6 Hz), 5.61 (1H, dd, *J* 0.6, 3.0 Hz), 5.12 (2H, s). LCMS (ES⁺) RT 3.418 minutes, 335 (M+H)⁺.

25

EXAMPLE 31-(4-Fluorobenzyl)-4-phenyl-1,4-dihydro-5H-pyrrolo[3,2-*b*]pyridin-5-one

δ_H (CDCl₃) 7.48 (2H, m), 7.37 (4H, m), 7.09 (2H, m), 7.00 (2H, m), 6.83 (1H, d, *J* 3.0 Hz), 6.37 (1H, d, *J* 9.5 Hz), 5.61 (1H, d, *J* 3.0 Hz), 5.14 (2H, s). LCMS (ES⁺) RT 3.252 minutes, 319 (M+H)⁺.

EXAMPLE 41-(2,6-Dichlorobenzyl)-4-phenyl-1,4-dihydro-5H-pyrrolo[3,2-b]pyridin-5-one

5 δ_H (CDCl₃) 7.61 (1H, d, *J* 9.5 Hz), 7.43 (2H, m), 7.32 (5H, m), 7.21 (1H, m), 6.65 (1H, d, *J* 3.1 Hz), 6.40 (1H, d, *J* 9.5 Hz), 5.49 (1H, d, *J* 3.1 Hz), 5.38 (2H, s). LCMS (ES⁺) RT 3.552 minutes, 369 (M+H)⁺.

EXAMPLE 510 1-(3-Methoxybenzyl)-4-phenyl-1,4-dihydro-5H-pyrrolo[3,2-b]pyridin-5-one

15 δ_H (CDCl₃) 7.48-7.46 (2H, m), 7.45-7.34 (4H, m), 7.23-7.19 (1H, m), 6.83 (1H, d, *J* 3.0 Hz), 6.78 (1H, dd, *J* 2.4, 8.3 Hz), 6.66 (1H, d, *J* 7.6 Hz), 6.60 (1H, s), 6.34 (1H, d, *J* 9.5 Hz), 5.58 (1H, d, *J* 3.0 Hz), 5.11(2H, s), 3.71 (3H, s). LCMS (ES⁺) RT 3.239 minutes, 331 (M+H)⁺.

15

EXAMPLE 61-Benzyl-4-(4-methoxyphenyl)-1,4-dihydro-5H-pyrrolo[3,2-b]pyridin-5-one

The title compound (9 mg) was prepared in a similar manner to the compound of Example 1 from Intermediate 7 (30 mg, 0.125 mmol) and benzyl chloride (18 mg, 0.14 mmol). δ_H (CDCl₃) 7.34 (1H, d, *J* 7.7 Hz), 7.28 (5H, m), 7.10 (2H, d, *J* 4.4 Hz), 6.98 (2H, d, *J* 6.6 Hz), 6.82 (1H, d, *J* 2.2 Hz), 6.33 (1H, d, *J* 7.7 Hz), 5.60 (1H, d, *J* 2.2 Hz), 5.15 (2H, s), 3.80 (3H, s). LCMS (ES⁺) RT 3.304 minutes, 331 (M+H)⁺.

25

EXAMPLE 71-Benzoyl-4-phenyl-1,4-dihydro-5H-pyrrolo[3,2-b]pyridin-5-one

The title compound (146 mg) was prepared, in a similar manner to Intermediate 4, from Intermediate 10. δ_H (d₆-DMSO) 8.32 (1H, d, *J* 9.8 Hz), 7.75 (3H, m), 7.59 (5H, m), 7.40 (2H, dd, *J* 1.5, 8.4 Hz), 7.30 (1H, d, *J* 3.7 Hz), 6.43 (1H, d, *J* 9.8 Hz), 5.81 (1H, dd, *J* 0.3, 4.1 Hz). LCMS (ES⁺) RT 3.299 minutes, 315 (M+H)⁺.

EXAMPLE 84-[5-Oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridin-1-yl)methyl]benzonitrile

From Intermediate 5 (150 mg, 0.71 mmol) and 4-cyanobenzyl bromide (139 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (151 mg, 65%). δ_H (d_6 -DMSO) 7.59 (2H, d, J 8.2 Hz), 7.50-7.45 (2H, m), 7.41-7.34 (3H, m), 7.26 (1H, d, J 9.1 Hz), 7.13 (2H, d, J 8.1 Hz), 6.83 (1H, d, J 3.0 Hz), 6.35 (1H, d, J 9.5 Hz), 5.64 (1H, d, J 2.7 Hz), 5.21 (2H, s). LCMS (ES $^+$) RT 3.018 minutes, 326 (M+H $^+$).

10

EXAMPLE 93-[5-Oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridin-1-yl)methyl]benzonitrile

From Intermediate 5 (150 mg, 0.71 mmol) and 3-cyanobenzyl bromide (139 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (148 mg, 64%). δ_H ($CDCl_3$) 7.55 (1H, d, J 7.7 Hz), 7.50-7.36 (6H, m), 7.33 (1H, s), 7.28 (2H, d, J 9.0 Hz), 6.83 (1H, d, J 3.0 Hz), 6.36 (1H, d, J 9.6 Hz), 5.65 (1H, dd, J 1.3, 2.4 Hz), 5.19 (2H, s). LCMS (ES $^+$) RT 3.021 minutes, 326 (M+H $^+$).

EXAMPLE 10

20

1-(2-Methylbenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 2-methylbenzyl bromide (131 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (129 mg, 58%). δ_H ($CDCl_3$) 7.59-7.54 (2H, m), 7.49-7.45 (4H, m), 7.30-7.19 (3H, m), 6.88 (1H, d, J 7.5 Hz), 6.79 (1H, d, J 3.0 Hz), 6.44 (1H, d, J 9.5 Hz), 5.66 (1H, d, J 3.0 Hz), 5.22 (2H, s), 2.33 (3H, s). LCMS (ES $^+$) RT 3.373 minutes, 315 (M+H $^+$).

EXAMPLE 111-(3-Methylbenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 3-methylbenzyl bromide (131 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (146 mg, 65%). δ_H ($CDCl_3$) 7.58-7.54 (2H, m), 7.48-7.44 (4H, m), 7.29-7.25 (1H, m), 7.15

(1H, d, *J* 7.6 Hz), 7.00-6.96 (2H, m), 6.91 (1H, d, *J* 3.0 Hz), 6.43 (1H, d, *J* 9.5 Hz), 5.67 (1H, d, *J* 3.0 Hz), 5.20 (2H, s), 2.37 (3H, s). LCMS (ES⁺) RT 3.363 minutes, 315 (M+H⁺).

5

EXAMPLE 121-(4-Methylbenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 4-methylbenzyl bromide (131 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (142 mg, 64%). δ_H (CDCl₃) 7.48-7.44 (2H, m), 7.39-7.34 (4H, m), 7.09 (2H, d, *J* 7.9 Hz), 6.98 (2H, d, *J* 8.2 Hz), 6.81 (1H, d, *J* 3.0 Hz), 6.33 (1H, d, *J* 9.4 Hz), 5.57 (1H, dd, *J* 0.6, 3.0 Hz), 5.10 (2H, s), 2.27 (3H, s). LCMS (ES⁺) RT 3.376 minutes, 315 (M+H⁺).

15

EXAMPLE 131-(4-Chlorobenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 4-chlorobenzyl bromide (146 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (158 mg, 67%). δ_H (CDCl₃) 7.57-7.54 (2H, m), 7.49-7.34 (6H, m), 7.09 (2H, d, *J* 8.3 Hz), 6.90 (1H, d, *J* 3.0 Hz), 6.43 (1H, d, *J* 9.5 Hz), 5.69 (1H, d, *J* 3.0 Hz), 5.21 (2H, s). LCMS (ES⁺) RT 3.383 minutes, 335 (M+H⁺).

EXAMPLE 1425 1-(3,4-Dichlorobenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 3,4-dichlorobenzyl bromide (170 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (173 mg, 66%). δ_H (CDCl₃) 7.59-7.55 (2H, m), 7.49-7.43 (5H, m), 7.41-7.26 (1H, m), 6.97 (1H, dd, *J* 2.1, 8.2 Hz), 6.90 (1H, d, *J* 3.0 Hz), 6.45 (1H, d, *J* 9.6 Hz), 5.71 (1H, d, *J* 3.0 Hz), 5.20 (2H, s). LCMS (ES⁺) RT 3.531 minutes, 369 (M+H⁺).

EXAMPLE 151-(2,5-Dichlorobenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 2,5-dichlorobenzyl bromide (170 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (147 mg, 40%). δ_H ($CDCl_3$) 7.62-7.52 (2H, m), 7.50-7.36 (5H, m), 7.30-7.27 (1H, m), 6.92 (1H, d, J 3.0 Hz), 6.78 (1H, d, J 3.4 Hz), 6.48 (1H, d, J 9.5 Hz), 5.74 (1H, d, J 3.0 Hz), 5.31 (2H, s). LCMS (ES $^+$) RT 3.525 minutes, 369 ($M+H^+$).

10

EXAMPLE 161-(3,4-Difluorobenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 3,4-difluorobenzyl bromide (147 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (154 mg, 64%). δ_H ($CDCl_3$) 7.59-7.54 (2H, m), 7.50-7.39 (4H, m), 7.22-7.15 (1H, m), 7.02-6.89 (3H, m), 6.45 (1H, d, J 9.5 Hz), 5.71 (1H, d, J 3.0 Hz), 5.20 (2H, s). LCMS (ES $^+$) RT 3.241 minutes, 337 ($M+H^+$).

20

EXAMPLE 171-(2,4-Difluorobenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 2,4-difluorobenzyl bromide (147 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (138 mg, 58%). δ_H ($CDCl_3$) 7.57-7.48 (3H, m), 7.47-7.41 (3H, m), 7.05-7.01 (1H, m), 6.99-6.86 (3H, m), 6.47 (1H, d, J 9.5 Hz), 5.68 (1H, d, J 2.9 Hz), 5.24 (2H, s). LCMS (ES $^+$) RT 3.257 minutes, 337 ($M+H^+$).

EXAMPLE 18

30 1-(3-Chloro-4-fluorobenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (150 mg, 0.71 mmol) and 3-chloro-4-fluorobenzyl bromide (159 mg, 0.71 mmol) according to the procedure for Example 6, to give the *title compound* (148 mg, 59%). δ_H (d_6 -DMSO) 7.68-7.64 (2H, m), 7.59-7.49 (4H, m), 7.33-

7.30 (1H, m), 7.25 (1H, t, *J* 8.6 Hz), 7.14-7.10 (1H, m), 7.00 (1H, d, *J* 3.0 Hz), 6.55 (1H, d, *J* 9.5 Hz), 5.80 (1H, dd, *J* 0.5, 3.0 Hz), 5.29 (2H, s). LCMS (ES⁺) RT 3.394 minutes, 353 (M+H⁺).

5

EXAMPLE 19

4-Phenyl-1-(pyridin-4-ylmethyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 5 (125 mg, 0.6 mmol) in THF (5 ml) was treated with sodium hydride (60% dispersion in mineral oil, 60 mg, 1.5 mmol), 4-chloromethylpyridine hydrochloride (98.5 mg, 0.6 mmol) and DMSO (1 ml). The reaction was stirred at r.t. for 2 h then warmed to 60°C for 30 minutes. After quenching with water the product was extracted into DCM, dried (Na₂SO₄) and concentrated *in vacuo*. Chromatography (10% MeOH in EtOAc, silica) gave the *title compound* (68 mg, 38%). δ_H (CDCl₃) 8.54 (2H, dd, *J* 1.6, 4.5 Hz), 7.50-7.46 (2H, m), 7.41-7.36 (3H, m), 7.27 (1H, d, *J* 9.6 Hz), 6.93 (2H, m), 6.84 (1H, d, *J* 3.0 Hz), 6.36 (1H, d, *J* 9.5 Hz), 5.65 (1H, d, *J* 3.0 Hz), 5.15 (2H, s). LCMS (ES⁺) RT 1.822 minutes, 302 (M+H⁺).

EXAMPLE 20

20 **4-Phenyl-1-(pyridin-3-ylmethyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one**

From Intermediate 5 (125 mg, 0.6 mmol), 3-chloromethylpyridine hydrochloride (98.5 mg, 0.6 mmol) and sodium hydride (60% in mineral oil, 60 mg, 1.5 mmol), by the method of Example 19, to give the *title compound* (57 mg, 32%). δ_H (CDCl₃) 8.62 (1H, dd, *J* 1.4, 4.7 Hz), 8.54 (1H, d, *J* 2.0 Hz), 7.58-7.54 (2H, m), 7.49-7.42 (5H, m), 7.34-7.30 (1H, m), 6.92 (1H, d, *J* 3.0 Hz), 6.46 (1H, d, *J* 9.6 Hz), 5.71 (1H, d, *J* 3.0 Hz), 5.27 (2H, s). LCMS (ES⁺) RT 2.220 minutes, 302 (M+H⁺).

EXAMPLE 21

30 **4-Phenyl-1-(1-phenylethyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one**

From Intermediate 5 (125 mg, 0.6 mmol), α-methylbenzyl bromide (111 mg, 0.6 mmol) and sodium hydride (60% in mineral oil, 31 mg, 0.78 mmol) by the method of Example 19 to give the *title compound* (76 mg, 40%). δ_H (CDCl₃) 7.58-7.54 (2H, m),

7.49-7.44 (3H, m), 7.41-7.31 (4H, m), 7.18-7.16 (2H, m), 7.08 (1H, d, *J* 3.1 Hz), 6.37 (1H, d, *J* 9.5 Hz), 5.70 (1H, d, *J* 3.1 Hz), 5.54 (1H, q, *J* 7.1 Hz), 1.92 (3H, d, *J* 7.06 Hz). LCMS (ES⁺) RT 3.320 minutes, 315 (M+H⁺).

5

EXAMPLE 221-(3-Chlorobenzyl)-4-phenyl-2-(pyrrolidin-1-ylsulfonyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

Intermediate 12 (140 mg, 0.408 mmol) in THF (10 ml) was treated with sodium hydride (60% dispersion in mineral oil, 19.5 mg, 0.49 mmol) under nitrogen at r.t. After stirring for 10 minutes 3-chlorobenzyl bromide (84.3 mg, 0.41 mmol) was added and the reaction stirred for 3 h. After 3 h TLC showed little change so further sodium hydride (60% dispersion in mineral oil, 20 mg, 0.41 mmol) was added followed by DMSO (1 ml), and the reaction heated at 70°C for 1 h then at r.t. overnight. The reaction was partitioned between water and DCM, and the organic phase separated, dried (Na₂SO₄), and concentrated *in vacuo*. Purification by chromatography (EtOAc-silica) gave the *title compound* (62 mg, 33%). δ_H (d₆-DMSO) 7.88-7.85 (1H, m), 7.63-7.58 (2H, m), 7.54-7.50 (1H, m), 7.47-7.44 (2H, m), 7.39-7.34 (2H, m), 7.22 (1H, s), 6.97-6.94 (1H, m), 6.48 (1H, d, *J* 9.8 Hz), 6.07 (1H, s), 5.71 (2H, s), 3.09-3.04 (4H, m), 1.65-1.60 (4H, m). LCMS (ES⁺) RT 3.283 minutes, 468 (M+H⁺).

EXAMPLE 23Ethyl 1-benzyl-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

Phenylboronic acid (47 mg, 0.37 mmol), pyridine (0.05 ml, 0.57 mmol) and copper(II) acetate (69 mg, 0.37 mmol) were added to a suspension of Intermediate 17 (57 mg, 0.19 mmol) in DCM (2 ml). The reaction mixture was stirred overnight at r.t. The reaction mixture was diluted with DCM (20 ml), washed with 2M HCl, dried (MgSO₄) and concentrated *in vacuo*. Purification by chromatography (10%-20% EtOAc in DCM) gave the *title compound* (44 mg, 62%). δ_H (CDCl₃) 7.20-7.60 (9H, m), 7.04-7.00 (2H, m), 6.51 (1H, d, *J* 9.7 Hz), 6.30 (1H, d, *J* 0.6 Hz), 5.71 (2H, s), 4.18 (2H, q, *J* 7.1 Hz), 1.20 (3H, t, *J* 7.1 Hz). LCMS Method C (ES⁺) RT 2.44 minutes, 373 (M+H⁺).

EXAMPLE 24Ethyl 1-(3-chloro-4-fluorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

5 From Intermediate 20 (880 mg, 2.53 mmol), phenylboronic acid (617 mg, 5.06 mmol), pyridine (0.61 ml, 7.59 mmol) and copper(II) acetate (920 mg, 5.06 mmol) according to the procedure for Example 23. Purification by chromatography (15% EtOAc in DCM, silica) gave the *title compound* (630 mg, 59%). δ_H ($CDCl_3$) 7.45-7.25 (6H, m), 7.03-6.76 (3H, m), 6.49 (1H, d, J 9.7 Hz), 6.24 (1H, s), 5.56 (2H, s), 4.11 (2H, q, J 7.1 Hz), 1.14 (3H, t, J 7.1 Hz). LCMS (ES $^+$) RT 3.83 minutes, 425 [^{35}Cl]:427 [^{37}Cl] 3:1 (M+H $^+$).

10

EXAMPLE 25Ethyl 1-(3-methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 23 (354 mg, 1.14 mmol), phenylboronic acid (278 mg, 2.28 mmol), pyridine (0.28 ml, 3.42 mmol) and copper(II) acetate (414 mg, 2.28 mmol) according to the procedure for Example 23. Purification by chromatography (15% EtOAc in DCM, silica) gave the *title compound* (348 mg, 79%). δ_H ($CDCl_3$) 7.57-7.34 (6H, m), 7.15-7.10 (1H, m), 7.00 (1H, d, J 7.6 Hz), 6.84 (1H, s), 6.78 (1H, d, J 7.6 Hz), 6.52 (1H, d, J 9.5 Hz), 6.30 (1H, s), 5.67 (2H, s), 4.18 (2H, q, J 7.1 Hz), 2.24 (3H, s), 1.20 (3H, t, J 7.1 Hz). LCMS (ES $^+$) RT 3.77 minutes, 387 (M+H $^+$).

25

EXAMPLE 26Ethyl 1-(3-chlorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 26 (300 mg, 0.91 mmol), phenylboronic acid (222 mg, 1.82 mmol), pyridine (0.22 ml, 2.73 mmol) and copper(II) acetate (330 mg, 1.82 mmol) according to the procedure for Example 23. Purification by chromatography (15% EtOAc in DCM, silica) gave the *title compound* (317 mg, 86%). δ_H ($CDCl_3$) 7.50-7.31 (6H, m), 7.16-7.14 (2H, m), 6.97 (1H, s), 6.87-6.85 (1H, m), 6.51 (1H, d, J 9.7 Hz), 6.28 (1H, s),

5.64 (2H, s), 4.15 (2H, q, *J* 7.1 Hz), 1.17 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.80 minutes, 407 [³⁵Cl]:409 [³⁷Cl] 3:1 (M+H⁺).

EXAMPLE 27

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1-(3-Chloro-4-fluorobenzyl)-N-methoxy-N-methyl-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

A suspension of Intermediate 27 (110 mg, 0.28 mmol) in DCM (2 ml) was treated with EDC (80 mg, 0.42 mmol), *N,O*-dimethylhydroxylamine (41 mg, 0.42 mmol) and triethylamine (0.12 ml, 0.84 mmol). The reaction was stirred at r.t. overnight. The reaction mixture was diluted with DCM (20 ml), washed with 2M HCl, dried (MgSO₄) and concentrated *in vacuo*. Purification by chromatography (20% EtOAc in DCM, silica) gave the *title compound* (36 mg, 29%). δ_H (CDCl₃) 7.48-7.34 (6H, m), 7.12-6.91 (3H, m), 6.52 (1H, d, *J* 9.7 Hz), 6.23 (1H, s), 5.52 (2H, s), 3.40 (3H, s), 3.18 (3H, s). LCMS (ES⁺) RT 3.37 minutes, 440 [³⁵Cl]:442 [³⁷Cl] 3:1 (M+H⁺).

EXAMPLE 28

N-Methoxy-*N*-methyl-1-(3-methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

From Intermediate 28 (29 mg, 0.08 mmol), using EDC (23 mg, 0.12 mmol), *N,O*-dimethylhydroxylamine (12 mg, 0.12 mmol) and triethylamine (0.034 ml, 0.24 mmol), according to the procedure of Example 27. Purification by chromatography (20% EtOAc in DCM, silica) gave the *title compound* (11 mg, 34%). δ_H (CDCl₃) 7.50-7.33 (6H, m), 7.19-7.11 (1H, m), 6.99 (1H, d, *J* 7.6 Hz), 6.87 (1H, s), 6.79 (1H, d, *J* 7.6 Hz), 6.48 (1H, d, *J* 9.7 Hz), 6.15 (1H, s), 5.52 (2H, s), 3.34 (3H, s), 3.15 (3H, s), 2.23 (3H, s). LCMS (ES⁺) RT 3.27 minutes, 402 (M+H⁺).

EXAMPLE 291-(3-Chlorobenzyl)-N-methoxy-N-methyl-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

5 From Intermediate 29 (56 mg, 0.15 mmol), using EDC (43 mg, 0.23 mmol), *N,O*-dimethylhydroxylamine (22 mg, 0.23 mmol) and triethylamine (0.06 ml, 0.45 mmol), according to the procedure of Example 27. Purification by chromatography (20% EtOAc in DCM, silica) gave the *title compound* (21 mg, 34%). δ_H ($CDCl_3$) 7.50-7.33 (5H, m), 7.19-7.16 (3H, m), 7.02 (1H, s), 6.94-6.91 (1H, m), 6.50 (1H, d, J 9.6 Hz), 6.21 (1H, s), 10 5.55 (2H, s), 3.38 (3H, s), 3.17 (3H, s). LCMS (ES^+) RT 3.33 minutes, 422 [^{35}Cl]:424 [^{37}Cl] 3:1 ($M+H^+$).

EXAMPLE 301-(3-Chloro-4-fluorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

A suspension of Intermediate 27 (100 mg, 0.25 mmol) in DMF (1 ml) was treated with CDI (50 mg, 0.30 mmol) and stirred at room temperature for 1 h. Ammonia 880 (1 ml) was added and the reaction mixture stirred at r.t. for 1 h. The reaction mixture was 20 partitioned between DCM and water (30 ml each) and the organics dried ($MgSO_4$) and concentrated *in vacuo*. Purification by chromatography (5% THF in EtOAc, silica) gave the *title compound* (65 mg, 65%). δ_H (d_6 -DMSO) 7.75 (1H, d, J 9.6 Hz), 7.69 (1H, br s), 7.38-7.01 (8H, m), 6.93-6.90 (1H, m), 6.19 (1H, s), 6.15 (1H, d, J 9.6 Hz), 5.63 (2H, s). LCMS (ES^+) RT 2.98 minutes, 396 [^{35}Cl]:398 [^{37}Cl] 3:1 ($M+H^+$).

25

EXAMPLE 311-(3-Methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

30 From Intermediate 28 (200 mg, 0.56 mmol) according to the procedure for Example 30. Purification by chromatography (5% THF in EtOAc, silica) gave the *title compound* (100 mg, 50%). δ_H (d_6 -DMSO) 7.94-7.87 (2H, m), 7.62-7.44 (5H, m), 7.30-

7.24 (2H, m), 7.12-7.07 (2H, m), 6.94 (1H, d, *J* 7.5 Hz), 6.43 (1H, s), 6.39 (1H, d, *J* 9.6 Hz), 5.90 (2H, s), 2.31 (3H, s). LCMS (ES⁺) RT 2.91 minutes, 358 (M+H⁺).

EXAMPLE 32

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1-(3-Chlorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

From Intermediate 29 (200 mg, 0.53 mmol) according to the procedure for Example 30. Purification by chromatography (5% THF in EtOAc, silica) gave the *title compound* (75 mg, 38%). δ_H (d₆-DMSO) 8.00-7.92 (2H, m), 7.67-7.53 (3H, m), 7.48-7.29 (6H, m), 7.16-7.11 (1H, m), 6.47 (1H, s), 6.42 (1H, d, *J* 9.6 Hz), 5.93 (2H, s). LCMS (ES⁺) RT 2.95 minutes, 378 [³⁵Cl]:380 [³⁷Cl] 3:1 (M+H⁺).

EXAMPLE 33

15

4-Phenyl-1-(2-phenylethyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

A solution of Intermediate 5 (210 mg, 1.00 mmol) in THF and DMF (5 ml each) was treated with sodium hydride (60% in mineral oil, 200 mg, 5.00 mmol) at r.t. 2-Phenylethyl bromide (0.68 ml, 5.00 mmol) was added and the reaction stirred at r.t. overnight. The reaction mixture was partitioned between water (30 ml) and EtOAc (40 ml), washed with brine (20 ml), dried (MgSO₄) and concentrated *in vacuo*. Purification by chromatography (50% EtOAc in DCM, silica) gave the *title compound* (218 mg, 68%). δ_H (CDCl₃) 7.82-7.52 (9H, m), 7.32-7.30 (2H, m), 6.94 (1H, d, *J* 2.8 Hz), 6.66 (1H, d, *J* 9.5 Hz), 5.79 (1H, d, *J* 2.8 Hz), 4.49 (2H, t, *J* 7.1 Hz), 3.32 (2H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.34 minutes, 315 (M+H⁺).

EXAMPLE 34

1-(3-Chlorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carbonitrile

30 A suspension of Example 32 (200 mg, 0.53 mmol) in DCM (2 ml) was treated with pyridine (0.11 ml, 1.06 mmol) followed, after a few minutes, by TFAA (0.09 ml, 0.64 mmol). The reaction was stirred at r.t. for 15 minutes. EtOH (2 ml) was added and stirred for 5 minutes prior to removal of solvents *in vacuo*. The residue was dissolved in

DCM (15 ml) and washed with copper(II) sulfate (2 x 10 ml), dried (MgSO_4) and concentrated *in vacuo* to give the *title compound* (175 mg, 92%). δ_{H} ($d_3\text{-MeOD}$) 7.87 (1H, d, J 9.7 Hz), 7.55-7.43 (3H, m), 7.32-7.23 (4H, m), 7.18 (1H, s), 7.06-7.03 (1H, m), 6.55 (1H, d, J 9.7 Hz), 6.25 (1H, s), 5.45 (2H, s). LCMS (ES^+) RT 3.43 minutes, 360
5 $[^{35}\text{Cl}]:[^{37}\text{Cl}]$ 3:1 ($\text{M}+\text{H}^+$).

EXAMPLE 35

1-(3-Chlorobenzyl)-N,N-dimethyl-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-
10 b]pyridine-2-carboxamide

A solution of Intermediate 29 (400 mg, 1.00 mmol) in DMF (5 ml) was treated sequentially with thionyl chloride (0.15 ml, 2.00 mmol), then 2M dimethylamine in THF (1.5 ml, 3.00 mmol). The reaction mixture was stirred at r.t. for 15 minutes. Water (20 ml) was added and the mixture extracted with DCM (2 x 20 ml), dried (MgSO_4),
15 concentrated *in vacuo* and purified by chromatography (EtOAc, silica) to give the *title compound* (200 mg, 50%). δ_{H} ($d_3\text{-MeOD}$) 7.86 (1H, d, J 9.5 Hz), 7.52-7.40 (3H, m), 7.33-7.31 (2H, m), 7.23-7.18 (2H, m), 7.10 (1H, s), 7.09-6.98 (1H, br m), 6.42 (1H, d, J 9.5 Hz), 5.77 (1H, s), 5.38 (2H, s), 2.95-2.83 (6H, br m). LCMS (ES^+) RT 3.12 minutes,
406 $[^{35}\text{Cl}]:[^{37}\text{Cl}]$ 3:1 ($\text{M}+\text{H}^+$).

20

EXAMPLE 36

1-(3-Chlorobenzyl)-N-methyl-5-oxo-4-phenyl-4,5-dihydro-1*H*-pyrrolo[3,2-b]pyridine-2-
carboxamide

25 From Intermediate 29 (400 mg, 1.00 mmol), using methylamine hydrochloride (203 mg, 3.00 mmol) and triethylamine (0.42 ml, 3.00 mmol), according to the procedure for Example 35. Purification by chromatography (EtOAc, silica) gave the *title compound* (104 mg, 27%). δ_{H} ($d_3\text{-MeOD}$) 7.92 (1H, dd, J 0.5, 9.5 Hz), 7.67-7.53 (3H, m), 7.46-7.43 (2H, m), 7.33-7.25 (2H, m), 7.17 (1H, s), 7.08-7.05 (1H, m), 6.53 (1H, d, J 9.6 Hz), 6.29
30 (1H, d, J 0.5 Hz), 5.88 (2H, s), 2.78 (3H, s). LCMS (ES^+) RT 3.12 minutes, 392
 $[^{35}\text{Cl}]:[^{37}\text{Cl}]$ 3:1 ($\text{M}+\text{H}^+$).

EXAMPLE 371-(3-Chlorobenzyl)-4-phenyl-2-(pyrrolidin-1-ylcarbonyl)-1,4-dihydro-5H-pyrrolo[3,2-b]pyridin-5-one

5 From Intermediate 29 (400 mg, 1.00 mmol), using pyrrolidine (0.25 ml, 3.00 mmol), following the procedure of Example 35. Purification by chromatography (EtOAc, silica) gave the *title compound* (102 mg, 24%). δ_H (d_3 -MeOD) 7.92 (1H, d, J , 9.6 Hz), 7.53-7.41 (3H, m), 7.34-7.31 (2H, m), 7.24-7.17 (2H, m), 7.07 (1H, s), 7.00-6.97 (1H, m), 6.44 (1H, d, J 9.6 Hz), 5.87 (1H, s), 5.45 (2H, s), 3.33 (2H, t, J 6.7 Hz), 3.13 (2H, t, J 6.7 Hz), 1.78-1.70 (2H, m), 1.64-1.57 (2H, m). LCMS (ES $^+$) RT 3.31 minutes, 432 [^{35}Cl]:434 [^{37}Cl] 3:1 (M $+\text{H}^+$).

10

EXAMPLE 38

15. 1-(3-Chlorobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-b]pyridine-2-carbohydrazide

A solution of Example 26 (200 mg, 0.49 mmol) in 2-ethoxyethanol (3 ml) was treated with hydrazine hydrate (0.10 ml, 2.04 mmol) and heated at 115°C overnight. The reaction mixture was concentrated *in vacuo*. Purification by chromatography (10%-50% EtOAc in DCM, silica) gave the *title compound* (60 mg, 31%). δ_H (d_3 -MeOD) 7.81 (1H, d, J 9.6 Hz), 7.55-7.41 (3H, m), 7.34-7.31 (2H, m), 7.21-7.13 (2H, m), 7.06 (1H, s), 6.99-6.95 (1H, m), 6.42 (1H, d, J 9.6 Hz), 6.17 (1H, s), 5.76 (2H, s). LCMS (ES $^+$) RT 2.78 minutes, 393 [^{35}Cl]:395 [^{37}Cl] 3:1 (M $+\text{H}^+$).

25

EXAMPLE 39Ethyl 1-(3-chlorobenzyl)-4-(1H-indol-5-yl)-5-oxo-4,5-dihydro-1H-pyrrolo[3,2-b]pyridine-2-carboxylate

From Intermediate 26 (500 mg, 1.51 mmol) and 5-indolylboronic acid (488 mg, 3.03 mmol) according to the procedure for Example 23. Purification by chromatography (10%-50% EtOAc in DCM, silica) gave the *title compound* (300 mg, 44%). δ_H (d_3 -MeOD) 7.96 (1H, d, J 9.8 Hz), 7.51-7.48 (2H, m), 7.43-7.41 (1H, m), 7.33-7.26 (2H, m), 7.17 (1H, s), 6.98-6.92 (2H, m), 6.46-6.41 (2H, m), 6.05 (1H, s), 5.76 (2H, s), 4.09 (2H, q,

J 7.1 Hz), 1.07 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.62 minutes, 446 [³⁵Cl]:448 [³⁷Cl] 3:1 (M+H⁺).

EXAMPLE 40

5

Ethyl 1-(3-chlorobenzyl)-5-oxo-4-(3-thienyl)-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

From Intermediate 26 (500 mg, 1.51 mmol) and thiophene-3-boronic acid (388 mg, 3.03 mmol) according to the procedure for Example 23. Purification by chromatography (10%-50% EtOAc in DCM, silica) gave the *title compound* (100 mg, 16%). δ_{H} (d_3 -MeOD) 7.44-7.38 (3H, m), 7.19-7.15 (3H, m), 7.00 (1H, s), 6.90-6.84 (1H, m), 6.53-6.41 (2H, m), 5.67 (2H, s), 4.21 (2H, q, *J* 7.1 Hz), 1.23 (3H, t, *J* 7.1 Hz). LCMS (ES⁺) RT 3.80 minutes, 413 [³⁵Cl]:415 [³⁷Cl] 3:1 (M+H⁺).

15

EXAMPLE 41

1-(3-Chlorobenzyl)-4-(1*H*-indol-5-yl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxamide

To a solution of Intermediate 30 (50 mg, 0.12 mmol) in DMF (1 ml) was added CDI (46 mg, 0.28 mmol) and the mixture stirred at r.t. for 1 h. The solution was treated with 880 ammonia (0.5 ml) and stirred for a further 15 min. The reaction mixture was partitioned between DCM and water (10 ml each) and neutralised with 2M HCl (pH 7-8). The mixture was extracted with DCM (2 x 10 ml), dried (MgSO_4) and concentrated *in vacuo*. Purification by chromatography (5% MeOH in DCM, silica) gave the *title compound* (15 mg, 30%). δ_{H} (d_3 -MeOD) 7.82 (1H, d, *J* 9.6 Hz), 7.51-7.48 (2H, m), 7.30-7.25 (3H, m), 7.07 (1H, s), 7.00-6.95 (2H, m), 6.47-6.44 (2H, m), 6.28 (1H, s), 5.80 (2H, s). LCMS (ES⁺) RT 2.95 minutes, 417 [³⁵Cl]:419 [³⁷Cl] 3:1 (M+H⁺).

30

EXAMPLE 42Ethyl 1-(4-fluoro-3-methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-b]pyridine-2-carboxylate

5 From Intermediate 33 (680 mg, 2.07 mmol) and phenylboronic acid (488 mg, 4.00 mmol) according to the procedure for Example 23. Purification by chromatography (5%-30% EtOAc in DCM, silica) gave the *title compound* (531 mg, 64%). δ_H ($CDCl_3$) 7.86 (1H, d, J 9.7 Hz), 7.56-7.44 (3H, s), 7.31 (2H, d, J 7.1 Hz), 6.95 (1H, d, J 7.2 Hz), 6.89-6.83 (2H, m), 6.48 (1H, d, J 9.7 Hz), 6.19 (1H, s), 5.71 (2H, s), 4.16 (2H, q, J 7.1 Hz),
10 2.11 (3H, s), 1.16 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 3.82 minutes, 405 ($M+H^+$).

EXAMPLE 431-(3-Chlorobenzyl)-4-(3-thienyl)-1,4-dihydro-5H-pyrrolo[3,2-b]pyridin-5-one

15 From Intermediate 35 (680 mg, 3.15 mmol) and 3-chlorobenzyl bromide (0.45 ml, 3.46 mmol) according to the procedure for Example 1. Purification by chromatography (10%-40% EtOAc in DCM, silica) gave the *title compound* (843 mg, 79%). δ_H ($CDCl_3$) 7.46-7.50 (2H, m), 7.40 (1H, d, J 9.5 Hz), 7.33-7.31 (2H, m), 7.26 (1H, dd, J 1.5, 4.9 Hz), 7.15 (1H, s), 7.04-7.00 (1H, m), 6.95 (1H, d, J 3.0 Hz), 6.42 (1H, d, J 9.5 Hz), 5.91 (1H, d, J 3.0 Hz), 5.22 (2H, s). LCMS (ES^+) RT 3.32 minutes, 341 [^{35}Cl]:343 [^{37}Cl] 3:1
20 ($M+H^+$).

EXAMPLE 44Ethyl 1-(3-chlorobenzyl)-4-(4-methylphenyl)-5-oxo-4,5-dihydro-1H-pyrrolo[3,2-b]pyridine-2-carboxylate

From Intermediate 26 (350 mg, 1.06 mmol) and 4-methylphenylboronic acid (288 mg, 2.12 mmol), according to the procedure for Example 23. Purification by chromatography (10-15% EtOAc in DCM, silica) gave the *title compound* (328 mg, 74%). δ_H ($CDCl_3$) 7.41 (1H, d, J 9.7 Hz), 7.32-7.18 (6H, s), 7.00 (1H, s), 6.90-6.88 (1H, m), 6.55 (1H, d, J 9.7 Hz), 6.33 (1H, s), 5.67 (2H, s), 4.18 (2H, q, J 7.1 Hz), 2.39 (3H, s), 1.21 (3H, t, J 7.1 Hz). LCMS (ES^+) RT 3.96 minutes, 421 [^{35}Cl]:423 [^{37}Cl] 3:1 ($M+H^+$).

EXAMPLE 451-(2-Cyanobenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxamide

5 From Intermediate 40 (314 mg, 0.85 mmol) using 880 ammonia (2 ml), according to the procedure for Example 35. Purification by chromatography (EtOAc, silica) gave the *title compound* (100 mg, 32%). δ_H (d_6 -DMSO) 7.75 (1H, d, *J* 9.7 Hz), 7.72-7.70 (2H, m), 7.47-7.24 (5H, m), 6.99 (1H, br s), 6.43-6.31 (2H, m), 6.21 (1H, d, *J* 9.7 Hz), 5.90 (2H, s). LCMS (ES⁺) RT 2.71 minutes, 369 (M+H⁺).

10

EXAMPLE 461-(4-Fluoro-3-methylbenzyl)-5-oxo-4-phenyl-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxamide

15 From Intermediate 41 (504 mg, 1.34 mmol) using 880 ammonia (2 ml), according to the procedure for Example 35. Purification by chromatography (EtOAc, silica) gave the *title compound* (220 mg, 44%). δ_H (d_6 -DMSO) 7.85 (1H, d, *J* 9.7 Hz), 7.59-7.40 (3H, m), 7.34-7.31 (2H, m), 7.08-7.03 (1H, m), 6.99 (1H, d, *J* 9.7 Hz), 6.92-6.87 (1H, m), 6.31 (1H, s), 6.29 (1H, d, *J* 9.7 Hz), 5.75 (2H, s), 2.11 (3H, s). LCMS (ES⁺) RT 2.95 minutes, 20 376 (M+H⁺).

EXAMPLE 471-(3-Chlorobenzyl)-4-(4-methylphenyl)-5-oxo-4,5-dihydro-1H-pyrrolo[3,2-*b*]pyridine-2-carboxamide

From Intermediate 42 (278 mg, 0.71 mmol) using 880 ammonia (2 ml), according to the procedure for Example 35. Purification by chromatography (20% EtOAc in DCM to neat EtOAc, silica) gave the *title compound* (168 mg, 74%). δ_H (d_6 -DMSO) 7.93-7.86 (1H, br m), 7.39-7.23 (6H, m), 7.19 (1H, s), 7.07-7.01 (1H, m), 6.39 (1H, s), 6.32 (1H, d, *J* 9.6 Hz), 5.84 (2H, s), 2.37 (3H, s). LCMS (ES⁺) RT 3.06 minutes, 392 [³⁵Cl]:394 [³⁷Cl] 3:1 (M+H⁺).

EXAMPLE 481-(3-Chlorobenzyl)-4-(4-methylphenyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carbohydrazide

5 From Example 44 (278 mg, 0.71 mmol) using hydrazine hydrate (3 ml), according to the procedure for Example 38. Purification by trituration with DCM gave the *title compound* (168 mg, 74%). δ_H (d_6 -DMSO) 7.96 (1H, d, J 9.7 Hz), 7.37-7.25 (6H, m), 7.19 (1H, s), 7.00 (1H, d, J 6.4 Hz), 6.44 (1H, s), 6.40 (1H, d, J 9.7 Hz), 5.82 (2H, s), 2.33 (3H, s). LCMS (ES $^+$) RT 2.93 minutes, 407 [^{35}Cl]:409 [^{37}Cl] 3:1 (M+H $^+$).

10

EXAMPLE 49Ethyl 1-(3-chlorobenzyl)-4-(2-nitrophenyl)-5-oxo-4,5-dihydro-1*H*-pyrrolo[3,2-*b*]pyridine-2-carboxylate

15 A suspension of Intermediate 26 (1.60 g, 4.85 mmol) in DMF (15 ml) was treated with sodium hydride (60% in mineral oil, 240 mg, 5.82 mmol) and stirred at r.t. for 20 mins. 2-Fluoronitrobenzene (1.02 ml, 9.70 mmol) was added and the reaction heated at 90°C for 3 days. The reaction mixture was concentrated *in vacuo*, partitioned between EtOAc (100 ml) and water (50 ml) and extracted with EtOAc (50 ml). The combined
20 organics were washed with brine, dried (MgSO_4) and concentrated *in vacuo*. Purification by chromatography (10%-15% EtOAc in DCM, silica) gave the *title compound* (120 mg, 5%). δ_H (CDCl_3) 8.17 (1H, d, J 8.0 Hz), 7.80-7.75 (1H, m), 7.65-7.60 (1H, m), 7.51 (1H, d, J 8.0 Hz), 7.47 (1H, d, J 9.8 Hz), 7.23-7.20 (2H, m), 7.03 (1H, s), 6.88-6.84 (1H, m), 6.50 (1H, d, J 9.8 Hz), 6.29 (1H, s), 5.85 (1H, d, J 16.2 Hz), 5.54 (1H, d, J 16.2 Hz), 4.19
25 (2H, q, J 7.0 Hz), 1.21 (3H, t, J 7.0 Hz). LCMS (ES $^+$) RT 3.68 minutes, 452 [^{35}Cl]:454 [^{37}Cl] 3:1 (M+H $^+$).

EXAMPLE 50

30 1-(1,3-Benzothiazol-2-ylmethyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (100 mg, 0.48 mmol) using 2-(bromomethyl)-1,3-benzothiazole (119 mg, 0.52 mmol) according to the procedure for Example 33.

Purification by chromatography (50% EtOAc in DCM to neat EtOAc, silica) gave the *title*

compound (80 mg, 50%). δ_H ($CDCl_3$) 7.98 (1H, d, J 8.7 Hz), 7.78 (1H, d, J 7.5 Hz), 7.54-7.34 (8H, m), 6.97 (1H, d, J 3.1 Hz), 6.40 (1H, d, J 9.5 Hz), 5.67 (1H, d, J 3.1 Hz), 5.54 (2H, s). LCMS (ES^+) RT 3.17 minutes, 358 ($M+H^+$).

5

EXAMPLE 511-[(5-Chloro-2-thienyl)methyl]-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 5 (100 mg, 0.48 mmol) using 2-chloro-5-(chloromethyl)-thiophene (0.063 ml, 0.52 mmol) according to the procedure for Example 33. Purification 10 by chromatography (50% EtOAc in DCM, silica) gave the *title compound* (150 mg, 92%). δ_H ($CDCl_3$) 7.50-7.32 (6H, m), 6.82 (1H, d, J 3.0 Hz), 6.73-6.68 (2H, m), 6.40 (1H, d, J 9.5 Hz), 5.58 (1H, d, J 3.0 Hz), 5.19 (2H, s). LCMS (ES^+) RT 3.35 minutes, 341 [^{35}Cl]:343 [^{37}Cl] 3:1 ($M+H^+$).

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EXAMPLE 521-Benzyl-4-phenyl-3-(trifluoroacetyl)-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

From Intermediate 45 (280 mg, 0.86 mmol) using copper(II) acetate (26 mg, 0.13 mmol), pyridine (0.25 ml, 2.50 mmol), pyridine *N*-oxide (238 mg, 2.50 mmol) and 20 phenylboronic acid (391 mg, 2.50 mmol) according to the procedure for Example 23. Purification by chromatography (EtOAc, silica) gave the *title compound* (77 mg, 22%). δ_H (d_6 -DMSO): 8.42-8.43 (1H, m), 7.93 (1H, d, J 9.7 Hz), 7.44-7.32 (8H, m), 7.24-7.21 (2H, m), 6.40 (1H, d, J 9.7 Hz), 5.58 (2H, s). LCMS (ES^+) RT 3.46 minutes, 397 ($M+H^+$).

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EXAMPLE 532-[(2*S*)-2-(Hydroxymethyl)pyrrolidin-1-yl]carbonyl}-1-(3-methylbenzyl)-4-phenyl-1,4-dihydro-5*H*-pyrrolo[3,2-*b*]pyridin-5-one

30 A suspension of Intermediate 28 (400 mg, 1.04 mmol) in DCM (20 ml) was treated with EDC (403 mg, 2.10 mmol) and HOBT (284 mg, 2.10 mmol) and stirred at r.t. for 1 h. (*S*)-Prolinol (0.21 ml, 2.10 mmol) was added and the reaction was stirred at r.t. overnight. The reaction mixture was diluted with DCM (20 ml), washed with water, dried

(Na₂SO₄) and concentrated *in vacuo*. Purification by chromatography (EtOAc, silica) gave the *title compound* (118 mg, 25%). δ_H (CDCl₃, 380K) 7.88 (1H, d, *J* 9.6 Hz), 7.56 (2H, m), 7.47 (1H, m), 7.39 (2H, m), 7.21 (1H, d, *J* 7.7 Hz), 7.09 (1H, m), 7.02 (1H, s), 6.96 (1H, d, *J* 7.7 Hz), 6.30 (1H, d, *J* 9.6 Hz), 5.84 (1H, s), 5.48 (1H, d, *J* 15.7 Hz), 5.42 5 (1H, d, *J* 15.7 Hz), 4.03 (1H, br s), 3.44-3.42 (1H, m), 3.37-3.36 (1H, m), 3.33-3.28 (2H, m), 2.29 (3H, s), 1.84-1.75 (3H, m), 1.57-1.55 (1H, m). LCMS (ES⁺) RT 2.40 minutes, 442 (M+H⁺).

BIOLOGICAL ASSAYS

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The following assays and animal models can be used to demonstrate the potency and selectivity of the compounds according to the invention. In each assay an IC₅₀ value was determined for each test compound and represents the concentration of compound necessary to achieve 50% inhibition.

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Preparation of activated human p38α for inhibitor assays

Purification of human p38α

Human p38α, incorporating an *N*-terminal (His)6 tag, was expressed in baculovirus-infected High-Five™ cells (Invitrogen) according to the manufacturer's 20 instructions. The cells were harvested 72 h post-infection and lysed in phosphate-buffered saline (PBS) containing 1% (w/v) β-octylglucoside and Complete, EDTA-free™ protease inhibitors (Roche Molecular Biochemicals). The lysate was centrifuged at 35000 x g for 30 min at 4°C and the supernatant applied to a NiNTA™ column (Qiagen). Bound protein was eluted by 150 mM imidazole in PBS (after a wash with 15 mM 25 imidazole in PBS) and directly applied to a HiTrap Q™ column (AP Biotech). Bound protein was eluted using a 20 column volume, 0 to 1 M NaCl gradient. Fractions containing (His)6-p38 were aliquoted and stored at -70°C prior to their activation.

Preparation of GST-MKK6EE-containing lysates

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E. coli (BL21 pLysS) expressing the constitutively-activated form of human MKK6 fused with an *N*-terminal glutathione-*S*-transferase tag (GST-MKK6EE) were harvested by centrifugation and frozen at -70°C. Cells were lysed by resuspension in 1/10th the culture volume of PBS containing Complete, EDTA-free™ protease inhibitors

followed by sonication on ice for 4 x 15 sec. Cell debris was removed by centrifugation at 35,000 x g and the resultant supernatant stored in aliquots at -70°C.

Activation of (His)6-p38

5 0.45 ml of purified (His)6-p38 was incubated with 50 µl of the GST-MKK6BE-containing lysate for 30 min at 23°C in the presence of 1 mM β-glycerophosphate, 10 mM MgCl₂ and 9 mM ATP. The extent of activation was monitored by mass spectrometric detection of the doubly-phosphorylated form of (His)6-p38, which routinely comprised greater than 90% of the final (His)6-p38 preparation. The activated (His)6-p38 was then
10 diluted x 10 in PBS and repurified using the method described above. The concentration of purified, activated (His)6-p38 was measured by UV absorbance at 280 nm using A₂₈₀, 0.1% = 1.2 and the preparation stored in aliquots at -70°C prior to its use in inhibitor assays.

15 p38 Inhibition Assays

Inhibition of phosphorylation of biotinylated myelin basic protein (MBP)

The inhibition of p38-catalysed phosphorylation of biotinylated MBP is measured using a DELFIA-based format. The assay was performed in a buffer comprising 20 mM HEPES (pH 7.4), 5 mM MgCl₂ and 3 mM DTT. For a typical IC₅₀ determination,
20 biotinylated MBP (2.5 µM) was incubated at room temperature in a streptavidin-coated microtitre plate together with activated gst-p38 (10 nM) and ATP (1 µM) in the presence of a range of inhibitor concentrations (final concentration of DMSO is 2 percent). After fifteen minutes the reaction was terminated by the addition of EDTA (75 mM). The microtitre plate was then washed with Tris-buffered saline (TBS), prior to the addition of
25 100 µl of anti-phospho MBP antibody (mouse) together with europium-labeled anti-mouse IgG antibody. After one hour at room temperature the plate was again washed in TBS followed by the addition of Enhancement solution (PerkinElmer Wallac). Fluorescence measurements were performed after a further fifteen minutes at room temperature. IC₅₀ values are determined from the plot of log₁₀[inhibitor concentration]
30 (x-axis) versus percentage inhibition of the fluorescence generated by a control sample in the absence of inhibitor (y-axis).

Purification of human Peripheral Blood Mononuclear Cells

Peripheral blood mononuclear cells (PBMC) were isolated from normal healthy volunteers. Whole blood was taken by venous puncture using heparinised vacutainers (Becton Dickinson), diluted 1 in 4 in RPMI 1640 (Gibco, UK) and centrifuged at 400 x g
5 for 35 min over a Ficoll-paque gradient (Amersham-Pharmacia Biotech, UK). Cells at the interface were removed and washed once followed by a low speed spin (250 x g) to remove platelets. Cells were then resuspended in DMEM containing 10% FCS, penicillin 100 units ml⁻¹, streptomycin 50 µg ml⁻¹ and glutamine 2 mM (Gibco, UK).

10 *Inhibitor dilutions*

Inhibitor stocks (20 mM) were kept as a frozen solution (-20°C) in DMSO. Serial dilutions of inhibitors were performed in DMSO as 250-times concentrated stocks. Inhibitors were diluted 1 in 250 into tissue culture media, prewarmed to 37°C and transferred to plates containing PBMC. PBMC and inhibitors were incubated together for
15 30 min prior to addition of LPS. Inhibitors used in whole blood assays were prepared according to a different regime. Using the same stock solution serial dilutions of inhibitors were performed in DMSO. Inhibitors were then diluted 1 in 500 straight into whole blood in a volume of 1 µl. Inhibitor was incubated with whole blood for 30 min prior to the addition of LPS.

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LPS stimulation of PBMC

PBMC were resuspended at a density of 2×10^5 cells/well in flat-bottomed 96-well tissue culture treated plates. After the addition of inhibitor cells were stimulated with an optimal dose of LPS (*E. coli* strain B5:055, Sigma, at a final concentration of 1
25 µgml⁻¹) and incubated at 37°C in 5% CO₂/95% air for 18 hours. TNF-α levels were measured from cell-free supernatants by sandwich ELISA (BioSource #CHC1751).

LPS stimulation of whole blood

Whole blood was taken by venous puncture using heparinised vacutainers (Becton
30 Dickinson), and 500 µl of blood aliquoted into each well of a 24-well tissue culture treated plate. After the addition of inhibitor cells were stimulated with an optimal dose of LPS (*E. coli* strain B5:055, Sigma, at a final concentration of 1 µgml⁻¹) and incubated at

37°C without CO₂ for 18 hours. TNF- α levels were measured from cell-free supernatants by sandwich BLISA (BioSource #CHC1751).

Rat LPS-induced TNF release

5 Male Lewis rats (180-200 g) are anaesthetised with Isofluor and injected i.v. with LPS* in a volume of 0.5 ml sterile saline. After 90 minutes blood is collected into EDTA tubes for preparation of plasma samples. Plasma is stored at -70°C prior to assay for TNF- α by commercial ELISA.

10 *Rat CIA*

Female Lewis rats (180-200 g) are anaesthetised with Isofluor and immunised i.d. at the base of the tail with 2 x 100 μ l of emulsion containing 4 mg/ml bovine collagen II in 0.01 M acetic acid and Freund's Incomplete Adjuvant at a ratio of 1:1. A polyarthritis develops with onset from about 13 days post-sensitisation. The disease is mainly

15 confined to the ankles and is quantified by plethysmometry. Results are expressed as change in paw volume over time.

Conclusion

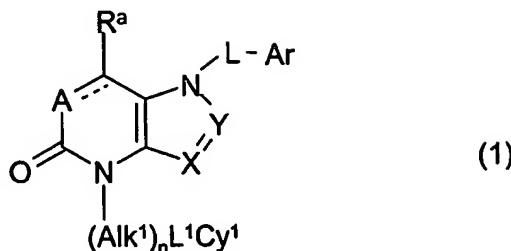
In the p38 inhibitor assays described above, the compounds of the Examples have

20 IC₅₀ values of around 2 μ M and below. The compounds of the invention are clearly potent inhibitors of p38 kinase, especially p38 α kinase.

Claims:

1. A compound of formula (1):

5



wherein:

- the dashed line joining A and C(R^a) is present and represents a bond and A is a -N= atom or a -C(R^b)= group, or the dashed line is absent and A is a -C(R^b)(R^c)- or
- 10 -N(R^d)- group;
- R^a, R^b and R^c is each independently a hydrogen or halogen atom or an optionally substituted alkyl, -CN, -CO₂R¹ (where R¹ is a hydrogen atom or an optionally substituted alkyl group) or -CONR¹R² group (where R² is a hydrogen atom or an optionally substituted alkyl group);
- 15 R^d is a hydrogen atom or an alkyl group;
- X and Y is each a nitrogen atom or a -C(R^e)= or -C(Alk²R^e)= group [where Alk² is an optionally substituted alkylene, alkenylene or alkynylene chain and R^e is a hydrogen or halogen atom or a -CN, -OR¹, -CO₂R¹, -C(X^a)R¹ (where X^a is an oxygen or sulphur atom), -Cy² (where Cy² is an optionally substituted, saturated or unsaturated non-aromatic carbocyclic ring optionally containing one or more -O-, -S-, -NH- or -C(X^a)- atoms or groups), -NR^{1a}R^{2a} (where R^{1a} and R^{2a}, which may be the same or different, is each a hydrogen atom or an optionally substituted alkyl or Cy² group, or together with the nitrogen atom to which they are attached form an optionally substituted, saturated or unsaturated cyclicamino ring optionally containing one or more -O- or -S- atoms or -NH- or -C(X^a)- groups), -C(X^a)NR^{1a}R^{2a}, -S(O)₂NR^{1a}R^{2a}, -N(R^{3a})C(X^a)R¹ (where R^{3a} is a hydrogen atom or an optionally substituted alkyl group), -N(R^{3a})C(X^a)NR^{1a}R^{2a}, -N(R^{3a})S(O)₂R¹, -N[S(O)₂R¹]₂, -N(R^{3a})S(O)₂NR^{1a}R^{2a}, -N(R^{3a})C(O)OR¹, -N(R^{3a})C(NR¹)NR^{1a}R^{2a}, -C(R¹)NOR², -C(NR¹)NR^{1a}R^{2a}, -C(X^a)NR^{1a}OR^{2a} or -C(O)N(R^{3a})NR^{1a}R^{2a} group];

L is a -C(O)-, -C(S)- or -C(R^{1f})(R^{1g})- group (where R^{1f} and R^{1g}, which may be the same or different, is each a hydrogen atom or a straight or branched C₁₋₃alkyl group optionally substituted by one, two or three fluorine atoms, or R^{1f} and R^{1g} together with the carbon atom to which they are attached form a cyclopropyl group), or a -CH₂CH₂- group;

5 n is zero or the integer 1;

Alk¹ is an optionally substituted aliphatic or heteroaliphatic chain;

L¹ is a covalent bond or a linker atom or group;

Cy¹ is an optionally substituted cycloaliphatic, polycycloaliphatic, heterocycloaliphatic, polyheterocycloaliphatic, aromatic or heteroaromatic group, or is

10 additionally a hydrogen atom when n is the integer 1 and/or L¹ is a linker atom or group; and

Ar is an optionally substituted aromatic or heteroaromatic group;

provided that the compound of formula (1) is other than 3,7-dibenzyl-3,7-dihydro-2H-purinone;

15 and the salts, solvates, hydrates and N-oxides thereof.

2. A compound as claimed in claim 1 wherein the dashed line joining A and C(R^a) is present and represents a bond and A is a -C(R^b)- group, in which R^a and R^b are as defined in claim 1.

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3. A compound as claimed in claim 2 wherein R^a and R^b are both hydrogen.

4. A compound as claimed in any one of the preceding claims wherein X is a -CH= group.

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5. A compound as claimed in any one of the preceding claims wherein Y is a -C(R^e)= group, in which R^e is hydrogen, -CN, -COR¹, -CO₂R¹, -CONR^{1a}R^{2a}, -S(O)₂NR^{1a}R^{2a}, -CONR^{1a}OR^{2a} or -C(O)N(R^{3a})NR^{1a}R^{2a}, and R¹, R^{1a}, R^{2a} and R^{3a} are as defined in claim 1.

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6. A compound as claimed in any one of the preceding claims wherein Cy¹ is phenyl, methylphenyl, methoxyphenyl, thienyl or indolyl.

7. A compound as claimed in any one of the preceding claims wherein Ar represents phenyl, fluorophenyl, difluorophenyl, chlorophenyl, dichlorophenyl, (chloro)(fluoro)phenyl, cyanophenyl, methylphenyl, (fluoro)(methyl)phenyl, methoxyphenyl, nitrophenyl, pyridinyl, chlorothienyl or benzothienyl.

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8. A compound as claimed in claim 1 as herein specifically disclosed in any one of the Examples.

9. A pharmaceutical composition comprising a compound of formula (1) as defined in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate or N-oxide thereof, in association with a pharmaceutically acceptable carrier.

10. The use of a compound of formula (1) as defined in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate or N-oxide thereof, for the manufacture of a medicament for the treatment and/or prevention of a disorder for which an inhibitor of p38 kinase is indicated.

11. A method for the treatment and/or prevention of a disorder for which an inhibitor of p38 kinase is indicated, which comprises administering to a patient in need of such treatment a compound of formula (1) as defined in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate or N-oxide thereof.

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB 03/04214

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07D471/04 A61K31/437 A61P19/00 A61P29/00 A61P37/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, BEILSTEIN Data, PAJ, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT
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Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, A	WO 03/033502 A (DAVIS JEREMY MARTIN ; BROOKINGS DANIEL CHRISTOPHER (GB); CELLTECH R &) 24 April 2003 (2003-04-24) claim 1	1-11
A	WO 99/64400 A (BEMIS GUY ; VERTEX PHARMA (US); COCHRAN JOHN (US); SALITURO FRANCESCO) 16 December 1999 (1999-12-16) claim 1	1-11

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

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Date of the actual completion of the International search	Date of mailing of the International search report
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26 January 2004

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Baston, E

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/GB 03/04214

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